WEST Search History

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DATE: Thursday, October 21, 2004

Hide?	Set Name	Query	Hit Count
	DB=PGPB,	USPT,USOC,EPAB,JPAB,DWPI; PLUR=Y	ES; OP=ADJ
	L12	ephrin-B3	14
	L11	L10 AND angiogenesis	19
	L10	AL-2 OR EFL-6 OR ephrin-B3	673
	L9	L6 AND ephrin-B3	0
	L8	L6 AND EFL-6	0
	L7	L6 AND AL-2	0
	L6	514/2.CCLS.	6340
	L5	Caras.IN.	179
	L4	Caras-I-W.IN.	8
	L3	Caras-I.IN.	1
	L2	Caras-Ingrid.IN.	2
	L1	(Caras-Ingrid-W.IN.)	16

END OF SEARCH HISTORY

Hit List

Clear Generate Collection Print Fwd Refs Bkwd Refs Generate OACS

Search Results - Record(s) 1 through 16 of 16 returned.

1. Document ID: US 20030049722 A1

Using default format because multiple data bases are involved.

L1: Entry 1 of 16

File: PGPB

Mar 13, 2003

PGPUB-DOCUMENT-NUMBER: 20030049722

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030049722 A1

TITLE: Novel methods of diagnosing macrophage developement related disorders, compositions, and methods of screening for macrophage developement modulators

PUBLICATION-DATE: March 13, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Murray, Richard	Cupertino	CA	US	
Caras, Ingrid W.	San Francisco	CA	US	
Hevezi, Peter	San Francisco	CA	US	
Wilson, Keith	Redwood City	CA	US	

US-CL-CURRENT: $\underline{435}/\underline{69.1}$; $\underline{435}/\underline{226}$, $\underline{435}/\underline{320.1}$, $\underline{435}/\underline{372}$, $\underline{435}/\underline{7.21}$, $\underline{536}/\underline{23.2}$

Full	Title Citation	Front Review	Classification	Date Referen	e Sequences	Attachments	Claims	KMC	Drawe Desi
			······						
	2. Docume	ent ID: US 20	020142444	4 1					
L1: E	ntry 2 of 1	16		File	: PGPB		Oc	t 3,	2002

PGPUB-DOCUMENT-NUMBER: 20020142444

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020142444 A1

TITLE: AL-2 neurotrophic factor

PUBLICATION-DATE: October 3, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Caras, Ingrid W. San Francisco CA US

US-CL-CURRENT: 435/226; 435/320.1, 435/325, 435/69.1, 536/23.2

ABSTRACT:

The present invention provides nucleic acids encoding AL-2 protein, as well as AL-2

h e b b g e e e f b e

protein produced by recombinant DNA methods. Such AL-2 protein and nucleic acid are useful in preparing antibodies and antagonists and in diagnosing and treating various neuronal disorders and disorders or conditions associated with angiogenesis.

Full Title Citation Front Review Classinostion Date Reference Sequences Attachments Claims Kill Diam Desi

3. Document ID: US 6696557 B1

L1: Entry 3 of 16

File: USPT

Feb 24, 2004

US-PAT-NO: 6696557

DOCUMENT-IDENTIFIER: US 6696557 B1

TITLE: AL-2 neurotrophic factor nucleic acid

DATE-ISSUED: February 24, 2004

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Caras; Ingrid W.

San Francisco

CA

US-CL-CURRENT: 536/23.4; 435/69.7

ABSTRACT:

The present invention provides nucleic acids encoding AL-2 protein, host cells and vectors containing these nucleic acids, and methods for their use to produce AL-2 protein by recombinant DNA methods.

2 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 15

Full Title Citation Front Review Classification Date Reference Citation Claims Killic Drams Desc

4. Document ID: US 6632634 B1

L1: Entry 4 of 16

File: USPT

CA

Oct 14, 2003

US-PAT-NO: 6632634

DOCUMENT-IDENTIFIER: US 6632634 B1

TITLE: Decay accelerating factor (DAF) and nucleic acid encoding it

DATE-ISSUED: October 14, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Caras; Ingrid W. San Francisco CA
Davitz; Michael A. Bronx NY
Nussenzweig; Victor New York NY

Martin, Jr.; David W. San Francisco

US-CL-CURRENT: 435/69.1; 435/252.3, 435/320.1, 435/325, 435/69.7, 536/23.5

ABSTRACT:

Novel fusions of a GPI signal domain and a polypeptide heterologous to the GPI signal domain donor polypeptide are provided for industrial use. Therapeutic administration of the GPI-linked product of the fusions enables the targeting of biological activity to cell membrane surfaces.

22 Claims, 19 Drawing figures Exemplary Claim Number: 1
Number of Drawing Sheets: 17

Fuil Title Citation Front Review Classification Date Reference Claims KMC Draw Desc

5. Document ID: US 6610296 B2

L1: Entry 5 of 16

File: USPT

Aug 26, 2003

US-PAT-NO: 6610296

DOCUMENT-IDENTIFIER: US 6610296 B2

TITLE: Methods of enhancing cognitive function using an AL-1 neurotrophic factor

immunoadhesin

DATE-ISSUED: August 26, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Caras; Ingrid W.

San Francisco

CA

Winslow; John W.

El Granada

CA

US-CL-CURRENT: 424/178.1; 514/12, 514/2, 530/350, 530/399

ABSTRACT:

The present invention provides methods of enhancing cognitive function in mammals by administering intracerebrally a homo-multimeric immunoadhesin molecule that contains the extracellular domain of AL-1, also known as ephrin-A5.

4 Claims, 7 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

Full Title Citation Front Review Classification Date Reference

6. Document ID: US 6280732 B1

L1: Entry 6 of 16

File: USPT

Aug 28, 2001

US-PAT-NO: 6280732

DOCUMENT-IDENTIFIER: US 6280732 B1

TITLE: Methods of using an AL-1 neurotrophic factor immunoadhesin

DATE-ISSUED: August 28, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Caras; Ingrid W.

San Francisco

CA

Winslow; John W.

El Granada

CA

US-CL-CURRENT: 424/178.1; 514/12, 514/2, 530/350, 530/399

ABSTRACT:

The present invention provides nucleic acids encoding AL-1 protein, as well as AL-1 protein produced by recombinant DNA methods. Such AL-1 protein is useful in preparing antibodies and in diagnosing and treating various neuronal disorders.

5 Claims, 7 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

Full	Title	Citation	Front	Review	Classification	Date	Reference		Claims	K966C	Draw Desi

7. Document ID: US 5798448 A

L1: Entry 7 of 16

File: USPT

Aug 25, 1998

US-PAT-NO: 5798448

DOCUMENT-IDENTIFIER: US 5798448 A

TITLE: AL-1 neurotrophic factor antibodies

DATE-ISSUED: August 25, 1998

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Caras; Ingrid W.

San Francisco

CA

Winslow; John W.

El Granada

CA

US-CL-CURRENT: 530/387.1; 424/130.1, 424/132.1, 424/133.1, 424/134.1, 424/135.1, 424/136.1, 424/9.34, 435/7.1, 435/7.2, 435/7.9, 436/512, 436/514, 436/517, 436/518, 436/536, 436/538, 436/547, 436/548, 530/300, 530/350, 530/387.3, 530/387.9, 530/388.1, 530/388.15, 530/388.24, 530/389.1

ABSTRACT:

The present invention provides nucleic acids encoding AL-1 protein, as well as AL-1 protein produced by recombinant DNA methods. Such AL-1 protein is useful in preparing antibodies and in diagnosing and treating various neuronal disorders.

9 Claims, 5 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 5

8. Document ID: US 5763224 A

L1: Entry 8 of 16

File: USPT

Jun 9, 1998

US-PAT-NO: 5763224

DOCUMENT-IDENTIFIER: US 5763224 A

** See image for Certificate of Correction **

TITLE: Decay accelerating factor (DAF) and nucleic acids encoding it

DATE-ISSUED: June 9, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Caras; Ingrid W. San Francisco CA Davitz; Michael A. Bronx NY Nussenzweig; Victor New York NY

Martin, Jr.; David W. San Francisco CA

US-CL-CURRENT: 435/69.6; 435/252.3, 435/320.1, 435/325, 435/455, 435/488, 435/69.7, <u>530/350, 530/829, 536/23.5</u>

ABSTRACT:

This application relates to nucleic acids encoding decay accelerating factor (hereinafter abbreviated as DAF), as well as vectors and cells which comprise such nucleic acids. Additionally, nucleic acids which encode variants of DAF, such as insertion, deletion or substitution variants, are described. This application also relates to the preparation of DAF in recombinant cell culture. In particular, it is concerned with the large scale manufacture of DAF suitable for pharmaceutical or diagnostic use.

33 Claims, 19 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 17

Full Title Citation Front	Berriem Classifica	tion Date Reference	2	Claims KilliGC Draint Desc
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9. Document ID: US 5759775 A

L1: Entry 9 of 16

File: USPT

Jun 2, 1998

US-PAT-NO: 5759775

DOCUMENT-IDENTIFIER: US 5759775 A

TITLE: Methods for detecting nucleic acids encoding AL--1 neurotrophic factor

DATE-ISSUED: June 2, 1998

INVENTOR-INFORMATION:

NAME STATE ZIP CODE COUNTRY CITY

Caras; Ingrid W. San Francisco CA

Winslow; John W. El Granada

e b g ee e f e b h ef h US-CL-CURRENT: 435/6; 435/91.2, 536/23.5, 536/24.31, 536/24.33

ABSTRACT:

Provided are nucleic acids encoding AL-1 protein, as well as AL-1 protein produced by recombinant DNA methods. Such AL-1 protein is useful in preparing antibodies and in diagnosing and treating various neuronal disorders. The present invention provides methods to preferentially detect or amplify AL-1 nucleic acid in a sample using AL-1 nucleotide sequence probes.

18 Claims, 7 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

Full	Title	Citation Front Review Classification Date Reference
	10.	Document ID: US 5374548 A

US-PAT-NO: 5374548

L1: Entry 10 of 16

DOCUMENT-IDENTIFIER: US 5374548 A

** See image for Certificate of Correction **

TITLE: Methods and compositions for the attachment of proteins to liposomes using a glycophospholipid anchor

DATE-ISSUED: December 20, 1994

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Nov 23, 1993

Dec 20, 1994

Caras; Ingrid W.

San Francisco

CA

File: USPT

US-CL-CURRENT: 424/450; 435/69.7, 436/829

ABSTRACT:

Novel fusions of a GPI signal domain and a polypeptide heterologous to the GPI signal domain donor polypeptide are provided for industrial use. Therapeutic administration of the GPI-linked product of the fusions enables the targeting of biological activity to cell membrane surfaces.

18 Claims, 19 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 19

Full	Title	Citation Front Review Classification Date Reference
	11.	Document ID: US 5264357 A

File: USPT

US-PAT-NO: 5264357

L1: Entry 11 of 16

DOCUMENT-IDENTIFIER: US 5264357 A

Record List Display Page 7 of 10

TITLE: Nucleic acids vectors and cells for the synthesis of membrane anchor fusion polypeptides

DATE-ISSUED: November 23, 1993

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Caras; Ingrid W. San Francisco CA
Davitz; Michael A. Riverdale NY
Nussenzweig; Victor New York NY
Martin, Jr.; David W. San Francisco CA

US-CL-CURRENT: <u>435/252.33</u>; <u>435/252.3</u>, <u>435/320.1</u>, <u>435/69.7</u>, <u>536/23.4</u>

ABSTRACT:

Novel fusions of a phospholipid anchor domain and a polypeptide heterologous to the anchor domain donor polypeptide are provided for industrial use. Therapeutic administration of the fusions enables the targeting of biological activity to cell membrane surfaces.

4 Claims, 14 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 14

Full Title Citation Front	Review Classification Da	te Reference	Claims KWIC Draw Desc

12. Document ID: US 5109113 A

L1: Entry 12 of 16

File: USPT

Apr 28, 1992

US-PAT-NO: 5109113

DOCUMENT-IDENTIFIER: US 5109113 A

TITLE: Membrane anchor fusion polypeptides

DATE-ISSUED: April 28, 1992

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Caras; Ingrid W. San Francisco CA
Davitz; Michael A. Riverdale NY
Nussenzweig; Victor New York NY
Martin, Jr.; David W. San Francisco CA

US-CL-CURRENT: 530/350; 435/69.7, 530/359, 530/405, 530/409, 530/806, 530/807, 530/808

ABSTRACT:

Novel fusions of a phospholipid anchor domain and a polypeptide heterologous to the anchor domain donor polypeptide are provided for industrial use. Therapeutic administration of the fusions enables the targeting of biological activity to cell membrane surfaces.

10 Claims, 14 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 14

Full Title Citation Front Review Classification Date Reference Claims Kint Dram Desi

13. Document ID: JP 08242882 A

L1: Entry 13 of 16

File: JPAB

Sep 24, 1996

PUB-NO: JP408242882A

DOCUMENT-IDENTIFIER: JP 08242882 A TITLE: NOVEL DAF AND ITS PRODUCTION

PUBN-DATE: September 24, 1996

INVENTOR-INFORMATION:

COUNTRY NAME

CARAS, INGRID W

INT-CL (IPC): $\underline{\text{C12}} \ \underline{\text{P}} \ \underline{\text{21/02}}; \ \underline{\text{C07}} \ \underline{\text{K}} \ \underline{\text{14/47}}; \ \underline{\text{C12}} \ \underline{\text{N}} \ \underline{\text{5/10}}; \ \underline{\text{C12}} \ \underline{\text{N}} \ \underline{\text{15/09}}$

ABSTRACT:

PROBLEM TO BE SOLVED: To provide a novel decay accelerating factor(DAF) that accelerates the decaying dissociation of C2 or the like from the C3 converting enzyme in the complement cascade that has a specific amino acid sequence and dissolves the antigen cells as the target of humoral immunity reaction and is useful for treatment and diagnosis of inflammatory diseases, autoimmune diseases and the like.

SOLUTION: This novel decay accelerating factor(DAF) has an amino acid sequence including the amino acid sequence represented by the formula and comprises the mDAF not accompanied by the natural glycosylation comprises the complement cascade dissolving the antigen cells as the target of the humoral immunity reaction.

COPYRIGHT: (C) 1996, JPO

Full Title Citation Front Review Classification Date Reference Claims KWIC Draw Desc

14. Document ID: WO 9740153 A1

L1: Entry 14 of 16

File: EPAB

Oct 30, 1997

PUB-NO: WO009740153A1

DOCUMENT-IDENTIFIER: WO 9740153 A1 TITLE: AL-2 NEUROTROPHIC FACTOR

PUBN-DATE: October 30, 1997

INVENTOR-INFORMATION:

NAME

COUNTRY

е

CARAS, INGRID W

INT-CL (IPC): <u>C12 N 15/12; C07 K 14/475; C12 N 15/62; A61 K 38/18; C07 K 16/22; C07 K</u> 19/00; C12 Q 1/68; G01 N 33/50; C12 N 5/10

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EUR-CL (EPC): C07K014/475

ABSTRACT:

CHG DATE=19990617 STATUS=O>The present invention provides nucleic acids encoding AL-2 protein, as well as AL-2 protein produced by recombinant DNA methods. Such AL-2 protein and nucleic acid are useful in preparing antibodies and antagonists and in diagnosing and treating various neuronal disorders and disorders or conditions associated with angiogenesis.

Full Title Citation Front Review Classification Date Reference Claims NWC Draws Described Descri

PUB-NO: WO009613518A1

DOCUMENT-IDENTIFIER: WO 9613518 A1

TITLE: AL-1 NEUROTROPHIC FACTOR, A LIGAND FOR AN EPH-RELATED TYROSINE KINASE RECEPTOR

PUBN-DATE: May 9, 1996

INVENTOR-INFORMATION:

NAME COUNTRY

CARAS, INGRID W US WINSLOW, JOHN W US

INT-CL (IPC): $\underline{\text{C07}}$ $\underline{\text{K}}$ $\underline{14}/\underline{47}$; $\underline{\text{C12}}$ $\underline{\text{N}}$ $\underline{15}/\underline{12}$; $\underline{\text{C07}}$ $\underline{\text{K}}$ $\underline{16}/\underline{18}$; $\underline{\text{C07}}$ $\underline{\text{K}}$ $\underline{19}/\underline{00}$; $\underline{\text{C12}}$ $\underline{\text{N}}$ $\underline{5}/\underline{10}$; $\underline{\text{C12}}$ $\underline{\text{Q}}$ $\underline{1/68}$; $\underline{\text{A61}}$ $\underline{\text{K}}$ $\underline{38}/\underline{12}$

EUR-CL (EPC): C07K016/22; C12N009/12, C07K014/52

ABSTRACT:

CHG DATE=19990617 STATUS=0>The present invention provides nucleic acids encoding AL-1 protein, as well as AL-1 protein produced by recombinant DNA methods. Such AL-1 protein is useful in preparing antibodies and antagonists and in diagnosing and treating various neuronal disorders and disorders or conditions associated with angiogenesis.

	Citation Front Review Classification			ims KMMC Draw Desc
	Document ID: WO 8901041 A1			······
L1: Entry	16 of 16	File:	EPAB	Feb 9, 1989

PUB-NO: WO008901041A1

DOCUMENT-IDENTIFIER: WO 8901041 A1

TITLE: NUCLEIC ACID AND METHODS FOR THE SYNTHESIS OF NOVEL FUSION POLYPEPTIDES WITH A

PHOSPHOLIPID ANCHOR DOMAIN

PUBN-DATE: February 9, 1989

INVENTOR-INFORMATION:

NAME

COUNTRY

CARAS, INGRID W

US

US-CL-CURRENT: 435/183; 435/320.1, 435/FOR.195, 530/387.3, 536/23.4

INT-CL (IPC): C12N 15/00

EUR-CL (EPC): G01N033/68; C07K014/705, C07K014/035

ABSTRACT:

CHG DATE=19990617 STATUS=O>Novel fusions of a phospholipid anchor domain and a polypeptide heterologous to the anchor domain donor polypeptide are provided for industrial use. Therapeutic administration of the fusions enables the targeting of biological activity to cell membrane surfaces.

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1. Document ID: US	20040096392 A1			
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L2: Entry 1 of 2	File:	PGPB		May 20, 2004
PGPUB-DOCUMENT-NUMBER: 200 PGPUB-FILING-TYPE: new DOCUMENT-IDENTIFIER: US 20				
TITLE: Antibodies against	cancer antigen TMEFF2	and uses t	hereof	
PUBLICATION-DATE: May 20,	2004			
INVENTOR-INFORMATION:				
NAME	CITY	STATE	COUNTRY	RULE-47
Bhaskar, Vinay	San Francisco	CA	US	
de la Calle, Agustin	Planegg	CA	DE	
Law, Debbie	San Francisco	CA	US	
Caras, Ingrid	San Francisco	CA	US	
Ramakrishnan, Vanitha	Belmont	CA	US	
Murray, Richard	Cupertino	CA	US	
Afar, Daniel	Fremont	CA	us	
Powers, David	Fairfax		us	
Full Title Coation Front Res D 2. Document ID: We L2: Entry 2 of 2	dem Classification Date Referen	ce Sequences		
PUB-NO: WO003075855A2 DOCUMENT-IDENTIFIER: WO 30 TITLE: ANTIBODIES AGAINST		AND USES T	THEREOF	
PUBN-DATE: September 18, 2	003			
INVENTOR-INFORMATION:				
NAME			COUNTRY	
BHASKAR, VINAY			US	
DE, LA CALLE AGUSTIN			DE	
LAW, DEBBIE			US	

US US

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CARAS, INGRID

RAMAKRISHNAN, VANITHA

MURRAY, RICHARD AFAR, DANIEL POWERS, DAVID US US US

INT-CL (IPC): $\underline{A61}$ \underline{K} $\underline{0}/$ EUR-CL (EPC): $\underline{C07K016/30}$

ABSTRACT:

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Search Results - Record(s) 1 through 1 of 1 returned.

1. Document ID: US 20040096392 A1, WO 2003075855 A2, AU 2003252830 A1 Using default format because multiple data bases are involved.

L3: Entry 1 of 1

File: DWPI

May 20, 2004

DERWENT-ACC-NO: 2003-756783

DERWENT-WEEK: 200434

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TITLE: New antibody that competitively inhibits binding of TMEFF219 to TMEFF2, useful for treating prostate cancer, e.g. primary, metastatic, locally advanced, or androgen

independent prostate cancer

INVENTOR: AFAR, D; BHASKAR, V; CARAS, I; DE LA CALLE, A; LAW, D; MURRAY, R;

POWERS, D ; RAMAKRISHNAN, V

PRIORITY-DATA: 2002US-436812P (December 27, 2002), 2002US-362837P (March 8, 2002),

2003US-0383447 (March 7, 2003)

PATENT-FAMILY:

PUB-NO PUB-DATE LANGUAGE PAGES MAIN-IPC US 20040096392 A1 May 20, 2004 000 A61K051/00 WO 2003075855 A2 September 18, 2003 031 A61K000/00 AU 2003252830 A1 September 22, 2003 000 A61K000/00

INT-CL (IPC): A61 \times 0/00; A61 \times 39/395; A61 \times 49/00; A61 \times 51/00

Full Ti	tle Citation	Front Review	Classification	Date Re	ference			Claims	KWC	Draw. Desc
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Search Results - Record(s) 1 through 8 of 8 returned.

1. Document ID: US 6632634 B1

Using default format because multiple data bases are involved.

L4: Entry 1 of 8

File: DWPI

Oct 14, 2003

DERWENT-ACC-NO: 2003-810556

DERWENT-WEEK: 200376

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TITLE: New nucleic acid comprising a sequence encoding a decay accelerating factor (DAF), useful for preparing a composition for treating inflammatory conditions, e.g., colitis, rheumatoid arthritis or allograft rejection

INVENTOR: CARAS, I W ; DAVITZ, M A ; MARTIN, D W ; NUSSENZWEIG, V

PRIORITY-DATA: 1993US-0017934 (February 12, 1993), 1985US-0738171 (May 24, 1985), 1986US-0859107 (May 1, 1986), 1987US-0083757 (August 6, 1987), 1991US-0811048 (December 19, 1991), 1994US-0358283 (December 19, 1994), 1998US-0014240 (January 27, 1998)

PATENT-FAMILY:

PUB-NO

PUB-DATE

LANGUAGE

PAGES :

MAIN-IPC

US 6632634 B1

October 14, 2003

035

C07H021/04

Mar 13, 2003

INT-CL (IPC): $\underline{\text{C07}} \ \underline{\text{H}} \ \underline{21/04}$

Full	Title	Citation Front Review Classification Date Reference Claims 1996 Draw Des
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	2.	Document ID: US 20030049722 A1

File: DWPI

DERWENT-ACC-NO: 2003-512353

DERWENT-WEEK: 200417

L4: Entry 2 of 8

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TITLE: Diagnosing destructive macrophage disorder (DMD) such as arthritis, aneurysms or atherosclerosis, or determining prognosis of individual with DMD, by determining expression or level of matrix metalloproteinase-19

INVENTOR: CARAS, I W ; HEVEZI, P ; MURRAY, R ; WILSON, K

PRIORITY-DATA: 2000US-0525978 (March 15, 2000), 1999US-124530P (March 15, 1999)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 US 20030049722 A1
 March 13, 2003
 061
 G01N033/53

h e b b g e e e f e b e f b

INT-CL (IPC): <u>C07 H 21/04</u>; <u>C12 N 5/08</u>; <u>C12 N 9/64</u>; <u>C12 P 21/02</u>; <u>G01 N 33/53</u>; <u>G01 N</u> 33/567

ABSTRACTED-PUB-NO: US20030049722A BASIC-ABSTRACT:

NOVELTY - Diagnosing (M1) destructive macrophage disorder (DMD) or determining (M2) prognosis of individual with DMD, by determining expression or level of matrix metalloproteinase (MMP)-19. (M1) involves comparing expression of gene encoding MMP-19 in a first tissue type of first individual, with expression of gene from second normal tissue from (I). (M2) involves determining level of MMP-19 in sample, where high level of MMP-19 indicates poor prognosis.

DETAILED DESCRIPTION - Diagnosing (M1) destructive macrophage disorder (DMD) or determining (M2) the prognosis of an individual with DMD, by determining the expression or level of MMP-19. (M1) involves determining expression of gene encoding or its fragment in a first tissue type of first individual (I), and comparing expression of the gene from second normal tissue from (I) or a second unaffected individual, where a difference in expression indicates that (I) has DMD. (M2) involves determining the level of MMP-19 in a sample, where a high level of MMP-19 indicates a poor prognosis.

INDEPENDENT CLAIMS are also included for the following:

- (1) screening (M3) drug candidates, involves providing a cell that expresses an expression profile gene which encodes a protein encoded by any one of 5 expression profile genes which are nucleic acids differentially expressed in the development path of destructive macrophages (DMs), referred as DM sequences, and the sequence represented by accession number X92521, X62466, J04130, X62078 and X76534, or its fragment; adding a drug candidate to the cell, and determining the effect of the drug candidate on the expression of the expression profile gene;
- (2) screening (M4) for a bioactive agent capable of binding to a destructive macrophage (DM) modulator protein or a bioactive agent capable of modulating the activity of a DM modulator protein, where the DM modulator protein is MMP-19 or its fragment, involves combining the DM modulator protein and a candidate bioactive agent, and determining the binding of the candidate agent to the DM modulator protein or determining the effect of the candidate agent on the bioactive of the DM modulator protein; evaluating (M5) the effect of a candidate DM drug, involves administering the drug to a patient, removing a cell sample from the patient, and determining the expression profile of the cell;
- (3) a biochip comprising a nucleic acid segment encoding MMP-19, or its fragment, where the biochip comprises fewer than 1000 nucleic acid probes;
- (4) an antibody (II) which specifically binds to MMP- 19, or its fragment;
- (5) screening for a bioactive agent capable of interfering with the binding of a DM modulator protein or its fragment and an antibody which binds to the DM modulator protein or its fragment, involves combining DM modulator protein or its fragment, a candidate bioactive agent and an antibody which binds to DM modulator protein or its fragment, and determining the binding of DM modulator protein or its fragment and the antibody;
- (6) inhibiting (M6) DMD, by administering to a cell a composition comprising (II);
- (7) inhibiting (M7) DMD in a cell by administering to a cell, a composition comprising antisense molecules to MMP- 19;
- (8) eliciting (M8) an immune response in an individual, by administering to the individual a composition comprising MMP-19 or its fragment, and optionally a carrier;
- (9) a composition (III) capable of eliciting an immune response in an individual,
- h eb bgeeef eb ef be

comprises nucleic acid encoding MMP-19 or its fragment and a carrier;

- (10) neutralizing the effect of a MMP-19 or its fragment by contacting an agent specific for the protein;
- (11) localizing (M9) a therapeutic moiety to colorectal cancer tissue by exposing the tissue to (II) conjugated to the therapeutic moiety; and
- (12) treating (M10) DMD by administering to the individual having DMD (II) conjugated to a therapeutic moiety.

ACTIVITY - Antiarthritic; Antiinflammatory; Antiatherosclerotic. No biological data is given.

MECHANISM OF ACTION - MMP-19-Inhibitor . No supporting data is given.

USE - (II) is useful for treating an individual for DMD by inhibiting MMP-19. (III) is useful for eliciting an immune response in an individual. (M6) is useful for inhibiting DMD in a cell of an individual having arthritis. (M8) is useful for eliciting an immune response in an individual. (M10) is useful for treating DMD (claimed). (II) is useful for treating a DMD such as arthritis, inflammatory bowel disease, chronic obstructive pulmonary disorder and vascular disease, including atherosclerosis and aneurysms. (II) is useful for inhibiting macrophage cell division, and for inhibiting macrophage development.

Full Title Citation

ef

Hit List

Clear Generate Collection	Print Fwd Rets	Bkwd Refs	Generate OACS

Search Results - Record(s) 1 through 19 of 19 returned.

1. Document ID: US 20040180823 A1

Using default format because multiple data bases are involved.

L11: Entry 1 of 19

File: PGPB

Sep 16, 2004

PGPUB-DOCUMENT-NUMBER: 20040180823

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040180823 A1

TITLE: Novel agents that modulate Eph receptor activity

PUBLICATION-DATE: September 16, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Pasquale, Elena B. San Diego CA US Koolpe, Mitchell San Diego CA US CA

Murai, Keith K. Candiac

US-CL-CURRENT: 514/12; 530/350

Full	Title Citation	Front F	Peview C	lassification	Date	Reference	Sequences	Attachments	Claims	KOME	Draw Desc

2. Document ID: US 20040136983 A1

L11: Entry 2 of 19 File: PGPB Jul 15, 2004

PGPUB-DOCUMENT-NUMBER: 20040136983

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040136983 A1

TITLE: Methods for inhibiting angiogenesis by EphB receptor antagonists

PUBLICATION-DATE: July 15, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Aguet, Michel Lutry CH

US-CL-CURRENT: 424/143.1

ABSTRACT:

The present application describes methods of inhibiting or stimulating angiogenesis in a mammal comprising administering to the mammal an effective amount of an Eph receptor antagonist or agonist, respectively. Articles of manufacture for use in

h e b b g ee e f e b ef b е relation to these methods are also described.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Affachinents | Claims | Kilifo | Draw Desc

3. Document ID: US 20040132634 A1

L11: Entry 3 of 19

File: PGPB

Jul 8, 2004

PGPUB-DOCUMENT-NUMBER: 20040132634

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040132634 A1

TITLE: Compositions and methods for regulating the kinase domain of receptor tyrosine

kinases

PUBLICATION-DATE: July 8, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Sicheri, Frank Toronto CA Wybenga-Groot, Leanne Etobicoke CA Pawson, Tony Toronto CA

US-CL-CURRENT: 514/1; 435/194, 702/19

ABSTRACT:

The present invention relates to binding pockets of receptor tyrosine kinases (RTKs). The binding pockets may regulate the kinase domain of the receptor tyrosine kinases. In particular, the invention relates to a crystal comprising a binding pocket of a receptor tyrosine kinase that regulates the kinase domain of the receptor tyrosine kinase EphB2. The crystal may be useful for modeling and/or synthesizing mimetics of a binding pocket or ligands that associate with the binding pocket. Such mimetics or ligands may be capable of acting as modulators of receptor tyrosine kinase receptor activity, and they may be useful for treating, inhibiting, or preventing diseases modulated by such receptors. Methods are also provided for regulating the kinase domain of an RTK by changing a binding pocket of the RTK that regulates the kinase domain from an autoinhibited state to an active state or from an active state to an autoinhibited state.

Full	Titie	Offation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KNINC	Draw, Desc

4. Document ID: US 20040126793 A1

L11: Entry 4 of 19

File: PGPB

Jul 1, 2004

PGPUB-DOCUMENT-NUMBER: 20040126793

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040126793 A1

TITLE: Lectin compositions and methods for modulating an immune response to an

antigen

PUBLICATION-DATE: July 1, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Segal, Andrew H. Boston MA US Young, Elihu Sharon MA US

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/419, 435/69.1, 530/370, 530/395,

<u>536/23.5</u>

ABSTRACT:

The present invention provides a fusion polypeptide which can bind to a cell surface binding moiety (e.g., a carbohydrate) and serve as a ligand for a cell surface polypeptide, as well as a vector comprising a nucleic acid encoding for such a fusion polypeptide, and a host cell comprising such nucleic acid. The present invention also provides a composition comprising an antigen bearing target and such a fusion polypeptide, as well as a composition comprising a virus or a cell and such a fusion polypeptide. The present invention further relates to a method of modulating an immune response in an animal using such compositions.

Full	Titie	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	K000C	Dram Desc
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5. Document ID: US 20040076955 A1

L11: Entry 5 of 19 File: PGPB Apr 22, 2004

PGPUB-DOCUMENT-NUMBER: 20040076955

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040076955 A1

TITLE: Methods of diagnosis of bladder cancer, compositions and methods of screening

for modulators of bladder cancer

PUBLICATION-DATE: April 22, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Mack, David H. Menlo Park CA US Aziz, Natasha Palo Alto CA US

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/69.1, 530/350, 536/23.5

ABSTRACT:

Described herein are genes whose expression are up-regulated or down-regulated in bladder cancer. Also described are such genes whose expression is further up-regulated or down-regulated in drug-resistant bladder cancer cells. Related methods and compositions that can be used for diagnosis, prognosis, or treatment of bladder cancer are disclosed. Also described herein are methods that can be used to identify modulators of bladder cancer.

Full Title Citation		Sequences Attachments Claims	
	 ·		

6. Document ID: US 20040002067 A1

L11: Entry 6 of 19

File: PGPB

Jan 1, 2004

PGPUB-DOCUMENT-NUMBER: 20040002067

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040002067 A1

TITLE: Breast cancer progression signatures

PUBLICATION-DATE: January 1, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Erlander, Mark G. Encinitas CA US Ma, Xia-Jun San Diego CA US

Sgroi, Dennis C. Winchester MA US

US-CL-CURRENT: 435/6; 435/287.2, 702/20

ABSTRACT:

Methods and compositions for the identification of breast cancer progression signatures are provided. The signature profiles are identified based upon multiple sampling of reference breast tissue samples from independent cases of breast cancer and provide a reliable set of molecular criteria for identification of cells as being in one or more particular stages of breast cancer.

Full			Classification			Claims	Drawi Desi

7. Document ID: US 20030157712 A1

L11: Entry 7 of 19 File: PGPB Aug 21, 2003

PGPUB-DOCUMENT-NUMBER: 20030157712

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030157712 A1

TITLE: Methods for determining cell responses through EphB receptors

PUBLICATION-DATE: August 21, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Daniel, Thomas O. Nashville TN US Stein, Elke San Francisco CA US

US-CL-CURRENT: 435/366; 435/368

ABSTRACT:

The present invention provides a method for initiating, promoting and/or directing cell attachment to a matrix or to another cell, comprising contacting an EphB receptor-expressing cell with a tetrameric EphB receptor-binding ligand, whereby binding of the tetrameric ligand promotes multimerization of the EphB receptor, thereby initiating, promoting and directing cell attachment to a matrix or to another cell. Also provided is a method for promoting <u>angiogenesis</u>, comprising contacting

Record List Display Page 5 of 13

EphB receptor-expressing cells which are associated with <u>angiogenesis</u> with a multimeric EphB receptor-binding ligand, whereby binding of the tetrameric ligand promotes multimerization of the EphB receptor, thereby promoting <u>angiogenesis</u>.

Full	Title	Citation Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KdaC	Dram. Desc
	**********				************				•	***********	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
	8.	Document ID:	US 200	30154032	Al						

File: PGPB

Aug 14, 2003

PGPUB-DOCUMENT-NUMBER: 20030154032

PGPUB-FILING-TYPE: new

L11: Entry 8 of 19

DOCUMENT-IDENTIFIER: US 20030154032 A1

TITLE: Methods and compositions for diagnosing and treating rheumatoid arthritis

PUBLICATION-DATE: August 14, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Pittman, Debra D.	Windham	NH	US	
Feldman, Jeffrey L.	Arlington	MA	US	
Shields, Kathleen M.	Harvard	MA	US	
Trepicchio, William L.	Andover	MA	US	

US-CL-CURRENT: 702/20

ABSTRACT:

The invention provides methods and compositions for diagnostic assays for detecting R.A. and therapeutic methods and compositions for treating R.A. The invention also provides methods for designing, identifying, and optimizing therapeutics for R.A. Diagnostic compositions of the invention include compositions comprising detection agents for detecting one or more genes that have been shown to be up- or down-regulated in cells of R.A. relative to normal counterpart cells. Exemplary detection agents include nucleic acid probes, which can be in solution or attached to a solid surface, e.g., in the form of a microarray. The invention also provides computer-readable media comprising values of levels of expression of one or more genes that are up- or down-regulated in R.A.

Full	Title Citation Front	Review Classification	Date Reference	Sequences	Attachments 0	laims Kee	C Draw De:
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	9. Document ID:	US 20030082511 A	<b>A</b> 1				
L11:	Entry 9 of 19	•	File:	PGPB		May 3	l, 2003

PGPUB-DOCUMENT-NUMBER: 20030082511

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030082511 A1

TITLE: Identification of modulatory molecules using inducible promoters

PUBLICATION-DATE: May 1, 2003

Jan 30, 2003

## Record List Display

INVENTOR-INFORMATION:

CITY COUNTRY RULE-47 NAME STATE Brown, Steven J. San Diego US CA Dunnington, Damien J. San Diego CA US Clark, Imran San Diego US CA

US-CL-CURRENT: 435/4; 435/6

#### ABSTRACT:

Methods for identifying an ion channel modulator, a target membrane receptor modulator molecule, and other modulatory molecules are disclosed, as well as cells and vectors for use in those methods. A polynucleotide encoding target is provided in a cell under control of an inducible promoter, and candidate modulatory molecules are contacted with the cell after induction of the promoter to ascertain whether a change in a measurable physiological parameter occurs as a result of the candidate modulatory molecule.

Full	Title	Citation Front Review	v Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw Desc
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	10.	Document ID: US	2003002220	2 A1						

File: PGPB

PGPUB-DOCUMENT-NUMBER: 20030022202

PGPUB-FILING-TYPE: new

L11: Entry 10 of 19

DOCUMENT-IDENTIFIER: US 20030022202 A1

TITLE: B-ephrin regulation of G-protein coupled chemoattraction, compositions, and methods of use

PUBLICATION-DATE: January 30, 2003

## INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Flanagan, John G. Newton US MA Brookline MA US Lu, Qiang Sun, Edna E. Brookline MA US

US-CL-CURRENT: 435/6; 435/196, 435/254.2, 435/320.1, 435/368, 435/69.1, 536/23.2

## ABSTRACT:

Transmembrane B ephrins and their Eph receptors signal bi-directionally. The presently claimed invention describes a cytoplasmic protein, designated PDZ-RGS3, which binds B ephrins through a PDZ domain, and has a regulator of heterotrimeric G protein signaling (RGS) domain. PDZ-RGS3 mediates signaling from the ephrin-B cytoplasmic tail. SDF-1, a chemokine with a G protein coupled receptor, or BDNF, act as chemoattractants for cerebellar granule cells, with SDF-1 action being selectively inhibited by soluble EphB receptor. The claimed invention reveals a pathway that links reverse signaling to cellular guidance, uncovers a novel mode of control for G proteins, and demonstrates a mechanism for selective regulation of responsiveness to neuronal guidance cues. Further, compositions and methods of use are provided for modulating cell migration as a function of chemokines and GPCR interaction, to aid in the treatment of disease states and medical conditions, including cancer and immune responses such as allergy and autoimmune responses. In one embodiment, a method of

altering the sensitivity of a cell to a chemokine is provided using a PDZ-RGS3 protein.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims Killic Draw Desc

11. Document ID: US 20020147306 A1

L11: Entry 11 of 19

File: PGPB

oct 10, 2002

PGPUB-DOCUMENT-NUMBER: 20020147306

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020147306 A1

TITLE: Peptides that modulate the interaction of B class ephrins and PDZ domains

PUBLICATION-DATE: October 10, 2002

INVENTOR-INFORMATION:

NAME CITY

STATE COUNTRY RULE-47

Lin, Danny Pawson, Anthony Scarborough Toronto CA CA

Gish, Gerald

East York

CA

US-CL-CURRENT: 530/350; 530/324

#### ABSTRACT:

The invention relates to complexes comprising a B class ephrin and a PDZ domain containing protein; peptides that interfere with the interaction of a B class ephrin with a PDZ domain binding site, and a PDZ domain containing protein; and, uses of the peptides and complexes. Methods for modulating the interaction of a B class ephrin and a PDZ domain containing protein, and methods for evaluating compounds for their ability to modulate the interaction are also described.

Fuil	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Drawt Desc
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12. Document ID: US 20020142444 A1

L11: Entry 12 of 19

File: PGPB

Oct 3, 2002

PGPUB-DOCUMENT-NUMBER: 20020142444

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020142444 A1

TITLE: AL-2 neurotrophic factor

PUBLICATION-DATE: October 3, 2002

INVENTOR-INFORMATION:

NAME CITY

STATE COUNTRY

RULE-47

Caras, Ingrid W.

San Francisco

CA

US

e

b

US-CL-CURRENT: 435/226; 435/320.1, 435/325, 435/69.1, 536/23.2

ABSTRACT:

The present invention provides nucleic acids encoding $\underline{AL-2}$ protein, as well as $\underline{AL-2}$ protein produced by recombinant DNA methods. Such $\underline{AL-2}$ protein and nucleic acid are useful in preparing antibodies and antagonists and in diagnosing and treating various neuronal disorders and disorders or conditions associated with <u>angiogenesis</u>.

Full Title Citation Front	Review Classification Date	Reference Sequences	Attachmento Claims KMC	Draw Desc
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13. Document ID: US 20020052308 A1

L11: Entry 13 of 19

File: PGPB

May 2, 2002

PGPUB-DOCUMENT-NUMBER: 20020052308

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020052308 A1

TITLE: Nucleic acids, proteins and antibodies

PUBLICATION-DATE: May 2, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Rosen, Craig A. Laytonsville MD US Ruben, Steven M. Olney MD US

US-CL-CURRENT: $\underline{514/1}$; $\underline{435/183}$, $\underline{435/320.1}$, $\underline{435/325}$, $\underline{435/6}$, $\underline{435/69.1}$, $\underline{435/7.1}$, $\underline{530/350}$,

<u>536/23.1</u>

ABSTRACT:

This invention relates to newly identified tissue specific cancer associated polynucleotides and the polypeptides encoded by these polynucleotides herein collectively known as "cancer antigens," and to the complete gene sequences associated therewith and to the expression products thereof, as well as the use of such tissue specific cancer antigens for detection, prevention and treatment of tissue specific disorders, particularly the presense of cancer. This invention relates to the cancer antigens as well as vectors, host cells, antibodies directed to cancer antigens and recombinant and synthetic methods for producing the same. Also provided are diagnostic methods for diagnosing and treating, preventing and/or prognosing tissue specific disorders, including cancer, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying agonists and antagonists of cancer antigens of the invention. The present invention further relates to methods and/or compositions for inhibiting the production and/or function of the polypeptides of the present invention.

Full Title Citation Front Review Classification Date Re	

14. Document ID: US 6727063 B1

L11: Entry 14 of 19

File: USPT

Apr 27, 2004

US-PAT-NO: 6727063

DOCUMENT-IDENTIFIER: US 6727063 B1

Feb 24, 2004

Record List Display

TITLE: Single nucleotide polymorphisms in genes

DATE-ISSUED: April 27, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Lander; Eric S. Cambridge MA Cargill; Michele Gaithersburg MD Ireland; James S. Gaithersburg West Roxbury Bolk; Stacey MA Daley; George Q. Weston McCarthy; Jeanette J. CA San Diego

US-CL-CURRENT: 435/6; 435/91.1, 435/91.2

ABSTRACT:

The invention provides nucleic acid segments of the human genome, particularly nucleic acid segments from a gene, including polymorphic sites. Allele-specific primers and probes hybridizing to regions flanking or containing these sites are also provided. The nucleic acids, primers and probes are used in applications such as phenotype correlations, forensics, paternity testing, medicine and genetic analysis. A role for the thrombospondin gene(s) in vascular disease is also disclosed. Use of single nucleotide polymorphisms in the thrombospondin gene(s) for diagnosis, prediction of clinical course and treatment response, development of therapeutics and development of cell-culture-based and animal models for research and treatment are disclosed.

4 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 8

Full Title Ci	lation Front Seview	Classification Date	Reference	Dlaims KOMC	Draw, Desc

15. Document ID: US 6696557 B1

L11: Entry 15 of 19 File: USPT

US-PAT-NO: 6696557

DOCUMENT-IDENTIFIER: US 6696557 B1

TITLE: AL-2 neurotrophic factor nucleic acid

DATE-ISSUED: February 24, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Caras; Ingrid W. San Francisco CA

US-CL-CURRENT: <u>536/23.4</u>; <u>435/69.7</u>

ABSTRACT:

The present invention provides nucleic acids encoding $\underline{AL-2}$ protein, host cells and vectors containing these nucleic acids, and methods for their use to produce $\underline{AL-2}$

protein by recombinant DNA methods.

2 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 15

Full Title Citation Front Review Classification Date Reference

16. Document ID: US 6555321 B1

L11: Entry 16 of 19

File: USPT

Apr 29, 2003

US-PAT-NO: 6555321

DOCUMENT-IDENTIFIER: US 6555321 B1

TITLE: Methods for determining cell responses through EphB receptors

DATE-ISSUED: April 29, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE COUNTRY

Daniel; Thomas O.

Nashville

TN

Stein; Elke

San Francisco

CA

US-CL-CURRENT: 435/7.1; 435/334, 435/7.2, 435/7.21, 435/7.8

ABSTRACT:

The present invention provides methods for screening an EphB receptor or an EphB receptor-binding ligand for the ability to promote a selected biological activity when in multimeric form. The invention also provides methods for initiating, promoting, directing, or inhibiting biological activities that involve EphB receptors and/or EphB receptor-binding ligands. The invention further provides compositions that can be used in the foregoing methods.

8 Claims, 9 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

> Full Title Citation Front Review Classification Date Reference Claims KMC Draw Des

17. Document ID: US 6514497 B1

L11: Entry 17 of 19

File: USPT

Feb 4, 2003

US-PAT-NO: 6514497

DOCUMENT-IDENTIFIER: US 6514497 B1

TITLE: Inhibition of LERK-2-mediated cell adhesion

DATE-ISSUED: February 4, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

e b b g ee e f e b ef h h e Briskin; Michael J.

Lexington Cambridge MA MA

US-CL-CURRENT: 424/143.1; 424/130.1, 424/137.1, 424/141.1, 424/152.1, 424/172.1, 530/387.1, 530/387.5, 530/388.1, 530/388.22

ABSTRACT:

Zou; Lily

Methods of modulating LERK-2-mediated cell adhesion, as well as methods of modulating <u>angiogenesis</u> and inflammation are described. Also described are agents such as antibodies which can modulate LERK-2-mediated cell adhesion, as well as methods of treating angiogenic diseases and inflammatory diseases.

8 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 8

Full	Citation	Review	Classification	Date	Reference		Claims	KMC	Draw Desc
	 _	 TT10		4					

18. Document ID: WO 9740153 A1

L11: Entry 18 of 19

File: EPAB

Oct 30, 1997

Feb 24, 2004

PUB-NO: WO009740153A1

DOCUMENT-IDENTIFIER: WO 9740153 A1 TITLE: AL-2 NEUROTROPHIC FACTOR

PUBN-DATE: October 30, 1997

INVENTOR-INFORMATION:

NAME

COUNTRY

CARAS, INGRID W

INT-CL (IPC): C12 N 15/12; C07 K 14/475; C12 N 15/62; A61 K 38/18; C07 K 16/22; C07 K

19/00; C12 Q 1/68; G01 N 33/50; C12 N 5/10

EUR-CL (EPC): C07K014/475

ABSTRACT:

CHG DATE=19990617 STATUS=0>The present invention provides nucleic acids encoding $\underline{AL-2}$ protein, as well as $\underline{AL-2}$ protein produced by recombinant DNA methods. Such $\underline{AL-2}$ protein and nucleic acid are useful in preparing antibodies and antagonists and in diagnosing and treating various neuronal disorders and disorders or conditions associated with $\underline{angiogenesis}$.

F	ull			Classification	Reference			Claims	KOME	Draw Desc
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19. Document ID: US 6696557 B1, WO 9740153 A1, AU 9726723 A, EP 904368 A1, AU 719273 B, JP 2000509978 W, US 20020142444 A1

File: DWPI

DERWENT-ACC-NO: 1997-535837

L11: Entry 19 of 19

DERWENT-WEEK: 200415

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TITLE: Human $\underline{AL-2}$ neurotrophic factor and related DNA - used to develop products for, e.g. treating neurologic disorders, $\underline{angiogenesis}$ disorders, tumours or rheumatoid arthritis or for wound healing

INVENTOR: CARAS, I W

PRIORITY-DATA: 1996US-0635130 (April 19, 1996), 2001US-0021121 (December 6, 2001)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
<u>US 6696557 B1</u>	February 24, 2004		000	C07H021/04
WO 9740153 A1	October 30, 1997	E	086	C12N015/12
<u>AU 9726723 A</u>	November 12, 1997		000	C12N015/12
EP 904368 A1	March 31, 1999	E	000	C12N015/12
<u>AU 719273 B</u>	May 4, 2000		000	C12N015/12
<u>JP 2000509978 W</u>	August 8, 2000		130	C12N015/09
US 20020142444 A1	October 3, 2002		000	C12N009/64

INT-CL (IPC): A61 K 38/18; C07 H 21/04; C07 K 14/475; C07 K 14/52; C07 K 16/22; C07 K 16/24; C07 K 19/00; C12 N 5/06; C12 N 5/10; C12 N 9/64; C12 N 15/09; C12 N 15/12; C12 N 15/62; C12 P 21/02; C12 P 21/08; C12 Q 1/68; G01 N 33/50

ABSTRACTED-PUB-NO: US20020142444A

BASIC-ABSTRACT:

A novel isolated nucleic acid (I) which encodes a polypeptide having an amino acid sequence that is at least 75% identical to 455 or 340 amino acid sequence for mature $\underline{AL-2}$ (given in the specification). Also claimed are: (1) an expression vector comprising (I) operably linked to a promoter; (2) a host cell transformed with an expression vector as in (1); (3) an isolated polypeptide as above; and (4) an antibody that specifically binds to a polypeptide as above.

USE - AL-2 is a novel Eph-related tyrosine kinase receptor ligand. AL-2 can be administered to patients in whom the nervous system has been damaged by trauma, surgery, stroke, ischaemia, infection, metabolic disease, nutritional deficiency, malignancy, or toxic agents, to promote the survival or growth of neurons. They can be used to treat motoneuron disorders such as amyotrophic lateral sclerosis (Lou Gehrig's disease), Bell's palsy, and various conditions involving spinal muscular atrophy, or paralysis. AL-2 can be used to treat human neurodegenerative disorders, such as Alzheimer's disease, Parkinson's disease, epilepsy, demyelinating diseases such as multiple sclerosis, Huntingtons chorea, Down's syndrome, nerve deafness, Menier's disease, and other disorders of the cerebellum. AL-2 can be used as cognitive enhancer, to enhance learning particularly in dementias or trauma, since they can promote axonal outgrowth and synaptic plasticity, particularly of hippocampal neurons that express AL-2 binding Eph-family receptors and cortical neurons that express AL-2. AL-2 can also be used for wound healing, i.e. accelerating neovascularisation of, e.g. burns and ulcers. AL-2 antagonists can be used for modulating angiogenesis. They can also be used for the treatment of tumours, acute myeloid leukaemia (AML), chronic myeloid leukaemia (CML), myelodysplastic syndrome (MDS), diabetic retinopathy, neovascular glaucoma, psoriasis and rheumatoid arthritis. The products can also be used for detection and diagnosis. ABSTRACTED-PUB-NO:

WO 9740153A EQUIVALENT-ABSTRACTS:

A novel isolated nucleic acid (I) which encodes a polypeptide having an amino acid sequence that is at least 75% identical to 455 or 340 amino acid sequence for mature

Record List Display Page 13 of 13

AL-2 (given in the specification). Also claimed are: (1) an expression vector comprising (I) operably linked to a promoter; (2) a host cell transformed with an expression vector as in (1); (3) an isolated polypeptide as above; and (4) an antibody that specifically binds to a polypeptide as above.

USE - AL-2 is a novel Eph-related tyrosine kinase receptor ligand. AL-2 can be administered to patients in whom the nervous system has been damaged by trauma, surgery, stroke, ischaemia, infection, metabolic disease, nutritional deficiency, malignancy, or toxic agents, to promote the survival or growth of neurons. They can be used to treat motoneuron disorders such as amyotrophic lateral sclerosis (Lou Gehrig's disease), Bell's palsy, and various conditions involving spinal muscular atrophy, or paralysis. AL-2 can be used to treat human neurodegenerative disorders, such as Alzheimer's disease, Parkinson's disease, epilepsy, demyelinating diseases such as multiple sclerosis, Huntingtons chorea, Down's syndrome, nerve deafness, Menier's disease, and other disorders of the cerebellum. AL-2 can be used as cognitive enhancer, to enhance learning particularly in dementias or trauma, since they can promote axonal outgrowth and synaptic plasticity, particularly of hippocampal neurons that express AL-2 binding Eph-family receptors and cortical neurons that express AL-2. AL-2 can also be used for wound healing, i.e. accelerating neovascularisation of, e.g. burns and ulcers. AL-2 antagonists can be used for modulating angiogenesis. They can also be used for the treatment of tumours, acute myeloid leukaemia (AML), chronic myeloid leukaemia (CML), myelodysplastic syndrome (MDS), diabetic retinopathy, neovascular glaucoma, psoriasis and rheumatoid arthritis. The products can also be used for detection and diagnosis.

Full	itle Citation	Front	Review	Classification	Date	Reference			Claim	s KOMO	Draw Des
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Search Results - Record(s) 1 through 14 of 14 returned.

1. Document ID: US 20040180823 A1

Using default format because multiple data bases are involved.

L12: Entry 1 of 14

File: PGPB

Sep 16, 2004

PGPUB-DOCUMENT-NUMBER: 20040180823

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040180823 A1

TITLE: Novel agents that modulate Eph receptor activity

PUBLICATION-DATE: September 16, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Pasquale, Elena B. San Diego CA US
Koolpe, Mitchell San Diego CA US
Murai, Keith K. Candiac CA

US-CL-CURRENT: <u>514/12</u>; <u>530/350</u>

	Citation Frent	ication Date	uences Attachments	Claims KMC	Draw, Desc
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2. Document ID: US 20040136983 A1

L12: Entry 2 of 14 File: PGPB Jul 15, 2004

PGPUB-DOCUMENT-NUMBER: 20040136983

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040136983 A1

TITLE: Methods for inhibiting angiogenesis by EphB receptor antagonists

PUBLICATION-DATE: July 15, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Aguet, Michel Lutry CH

US-CL-CURRENT: <u>424/143.1</u>

ABSTRACT:

The present application describes methods of inhibiting or stimulating angiogenesis in a mammal comprising administering to the mammal an effective amount of an Eph receptor antagonist or agonist, respectively. Articles of manufacture for use in

relation to these methods are also described.

Fuil	Title Citation Front	Review Classification Date	Reference S	Sequences	Attachmenta Clair	ns Kwic	Draw Desc
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	3. Document ID:	US 20040132634 A1					
L12:	Entry 3 of 14		File: P	PGPB		Jul 8,	2004

PGPUB-DOCUMENT-NUMBER: 20040132634

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040132634 A1

TITLE: Compositions and methods for regulating the kinase domain of receptor tyrosine

kinases

PUBLICATION-DATE: July 8, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47
Sicheri, Frank Toronto CA
Wybenga-Groot, Leanne Etobicoke CA
Pawson, Tony Toronto CA

US-CL-CURRENT: <u>514/1</u>; <u>435/194</u>, 702/19

ABSTRACT:

The present invention relates to binding pockets of receptor tyrosine kinases (RTKs). The binding pockets may regulate the kinase domain of the receptor tyrosine kinases. In particular, the invention relates to a crystal comprising a binding pocket of a receptor tyrosine kinase that regulates the kinase domain of the receptor tyrosine kinase EphB2. The crystal may be useful for modeling and/or synthesizing mimetics of a binding pocket or ligands that associate with the binding pocket. Such mimetics or ligands may be capable of acting as modulators of receptor tyrosine kinase receptor activity, and they may be useful for treating, inhibiting, or preventing diseases modulated by such receptors. Methods are also provided for regulating the kinase domain of an RTK by changing a binding pocket of the RTK that regulates the kinase domain from an autoinhibited state to an active state or from an active state to an autoinhibited state.

Full Title Citation Front Review Classification	Date Reference Sequences Atlach	ments Claims KMC Draw Desc
☐ 4. Document ID: US 2004012679		***************************************
L12: Entry 4 of 14	File: PGPB	Jul 1, 2004

PGPUB-DOCUMENT-NUMBER: 20040126793

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040126793 A1

TITLE: Lectin compositions and methods for modulating an immune response to an antigen

ancz 90-

PUBLICATION-DATE: July 1, 2004

Apr 22, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Segal, Andrew H. Boston MA US Young, Elihu Sharon MA US

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/419, 435/69.1, 530/370, 530/395,

<u>536/23.5</u>

ABSTRACT:

The present invention provides a fusion polypeptide which can bind to a cell surface binding moiety (e.g., a carbohydrate) and serve as a ligand for a cell surface polypeptide, as well as a vector comprising a nucleic acid encoding for such a fusion polypeptide, and a host cell comprising such nucleic acid. The present invention also provides a composition comprising an antigen bearing target and such a fusion polypeptide, as well as a composition comprising a virus or a cell and such a fusion polypeptide. The present invention further relates to a method of modulating an immune response in an animal using such compositions.

Full Title Citation Front Sevi	and Classification Date Reference	Sequences Atlachments Claims	KNOOC Drawn Desc
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File: PGPB

5. Document ID: US 20040076955 A1

PGPUB-DOCUMENT-NUMBER: 20040076955

PGPUB-FILING-TYPE: new

L12: Entry 5 of 14

DOCUMENT-IDENTIFIER: US 20040076955 A1

TITLE: Methods of diagnosis of bladder cancer, compositions and methods of screening

for modulators of bladder cancer

PUBLICATION-DATE: April 22, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Mack, David H. Menlo Park CA US Aziz, Natasha Palo Alto CA US

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/69.1, 530/350, 536/23.5

ABSTRACT:

Described herein are genes whose expression are up-regulated or down-regulated in bladder cancer. Also described are such genes whose expression is further up-regulated or down-regulated in drug-resistant bladder cancer cells. Related methods and compositions that can be used for diagnosis, prognosis, or treatment of bladder cancer are disclosed. Also described herein are methods that can be used to identify modulators of bladder cancer.

Full	Title	Citation	Fient	Review	Classification	Date	Reference	Sequences	Attachments Claims	K00C Draw Desc

6. Document ID: US 20040002067 A1

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L12: Entry 6 of 14

File: PGPB

Jan 1, 2004

PGPUB-DOCUMENT-NUMBER: 20040002067

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040002067 A1

TITLE: Breast cancer progression signatures

PUBLICATION-DATE: January 1, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Erlander, Mark G. Encinitas CA US
Ma, Xia-Jun San Diego CA US
Sgroi, Dennis C. Winchester MA US

US-CL-CURRENT: 435/6; 435/287.2, 702/20

ABSTRACT:

Methods and compositions for the identification of breast cancer progression signatures are provided. The signature profiles are identified based upon multiple sampling of reference breast tissue samples from independent cases of breast cancer and provide a reliable set of molecular criteria for identification of cells as being in one or more particular stages of breast cancer.

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Fu	#	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KOMBC	Drawa Desi

7. Document ID: US 20030157712 A1

File: PGPB Aug 21, 2003

PGPUB-DOCUMENT-NUMBER: 20030157712

PGPUB-FILING-TYPE: new

L12: Entry 7 of 14

DOCUMENT-IDENTIFIER: US 20030157712 A1

TITLE: Methods for determining cell responses through EphB receptors

PUBLICATION-DATE: August 21, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Daniel, Thomas O. Nashville TN US Stein, Elke San Francisco CA US

US-CL-CURRENT: 435/366; 435/368

ABSTRACT:

The present invention provides a method for initiating, promoting and/or directing cell attachment to a matrix or to another cell, comprising contacting an EphB receptor-expressing cell with a tetrameric EphB receptor-binding ligand, whereby binding of the tetrameric ligand promotes multimerization of the EphB receptor, thereby initiating, promoting and directing cell attachment to a matrix or to another cell. Also provided is a method for promoting angiogenesis, comprising contacting

EphB receptor-expressing cells which are associated with angiogenesis with a multimeric EphB receptor-binding ligand, whereby binding of the tetrameric ligand promotes multimerization of the EphB receptor, thereby promoting angiogenesis.

Full	Title	: Citation Fro	nt Review	Classification	Date	Reference	Sequences	Attachments 01	aims KodC	Draw Desi
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8. Document ID: US 20030154032 A1

L12: Entry 8 of 14

File: PGPB

Aug 14, 2003

PGPUB-DOCUMENT-NUMBER: 20030154032

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030154032 A1

TITLE: Methods and compositions for diagnosing and treating rheumatoid arthritis

PUBLICATION-DATE: August 14, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Pittman, Debra D.	Windham	NH	US	
Feldman, Jeffrey L.	Arlington	MA	US	
Shields, Kathleen M.	Harvard	MA	US	
Trepicchio, William L.	Andover	MA	US	

US-CL-CURRENT: 702/20

ABSTRACT:

The invention provides methods and compositions for diagnostic assays for detecting R.A. and therapeutic methods and compositions for treating R.A. The invention also provides methods for designing, identifying, and optimizing therapeutics for R.A. Diagnostic compositions of the invention include compositions comprising detection agents for detecting one or more genes that have been shown to be up- or down-regulated in cells of R.A. relative to normal counterpart cells. Exemplary detection agents include nucleic acid probes, which can be in solution or attached to a solid surface, e.g., in the form of a microarray. The invention also provides computer-readable media comprising values of levels of expression of one or more genes that are up- or down-regulated in R.A.

Full Title Cdation Front Serview	Classification Date Reference Sequences Attachments Claims)	GMC Draw Desc

9. Document ID: US 20030082511 A1

L12: Entry 9 of 14

File: PGPB

May 1, 2003

PGPUB-DOCUMENT-NUMBER: 20030082511

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030082511 A1

TITLE: Identification of modulatory molecules using inducible promoters

PUBLICATION-DATE: May 1, 2003

RULE-47

Jan 30, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY Brown, Steven J. San Diego CA US Dunnington, Damien J. San Diego CA US Clark, Imran San Diego CA US

US-CL-CURRENT: 435/4; 435/6

ABSTRACT:

Methods for identifying an ion channel modulator, a target membrane receptor modulator molecule, and other modulatory molecules are disclosed, as well as cells and vectors for use in those methods. A polynucleotide encoding target is provided in a cell under control of an inducible promoter, and candidate modulatory molecules are contacted with the cell after induction of the promoter to ascertain whether a change in a measurable physiological parameter occurs as a result of the candidate modulatory molecule.

Full Title C	itation Front Review Classification Date	Reference Sequences	Attachmenta Claims	KWWC Draw Desc
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L12: Entry	10 of 14	File: PGPB	.Tar	30 2002

File: PGPB

PGPUB-DOCUMENT-NUMBER: 20030022202

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030022202 A1

TITLE: B-ephrin regulation of G-protein coupled chemoattraction, compositions, and methods of use

PUBLICATION-DATE: January 30, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Flanagan, John G. Newton MA US Lu, Qiang Brookline MΑ US Sun, Edna E. Brookline US

US-CL-CURRENT: 435/6; 435/196, 435/254.2, 435/320.1, 435/368, 435/69.1, 536/23.2

ABSTRACT:

Transmembrane B ephrins and their Eph receptors signal bi-directionally. The presently claimed invention describes a cytoplasmic protein, designated PDZ-RGS3, which binds B ephrins through a PDZ domain, and has a regulator of heterotrimeric G protein signaling (RGS) domain. PDZ-RGS3 mediates signaling from the ephrin-B cytoplasmic tail. SDF-1, a chemokine with a G protein coupled receptor, or BDNF, act as chemoattractants for cerebellar granule cells, with SDF-1 action being selectively inhibited by soluble EphB receptor. The claimed invention reveals a pathway that links reverse signaling to cellular guidance, uncovers a novel mode of control for G proteins, and demonstrates a mechanism for selective regulation of responsiveness to neuronal guidance cues. Further, compositions and methods of use are provided for modulating cell migration as a function of chemokines and GPCR interaction, to aid in the treatment of disease states and medical conditions, including cancer and immune responses such as allergy and autoimmune responses. In one embodiment, a method of

e

May 2, 2002

altering the sensitivity of a cell to a chemokine is provided using a PDZ-RGS3 protein.

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KodC Drawt Des-

File: PGPB

PGPUB-DOCUMENT-NUMBER: 20020052308

PGPUB-FILING-TYPE: new

L12: Entry 11 of 14

DOCUMENT-IDENTIFIER: US 20020052308 A1

TITLE: Nucleic acids, proteins and antibodies

PUBLICATION-DATE: May 2, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Rosen, Craig A. Laytonsville MD US
Ruben, Steven M. Olney MD US

US-CL-CURRENT: 514/1; 435/183, 435/320.1, 435/325, 435/6, 435/69.1, 435/7.1, 530/350, 536/23.1

ABSTRACT:

This invention relates to newly identified tissue specific cancer associated polynucleotides and the polypeptides encoded by these polynucleotides herein collectively known as "cancer antigens," and to the complete gene sequences associated therewith and to the expression products thereof, as well as the use of such tissue specific cancer antigens for detection, prevention and treatment of tissue specific disorders, particularly the presense of cancer. This invention relates to the cancer antigens as well as vectors, host cells, antibodies directed to cancer antigens and recombinant and synthetic methods for producing the same. Also provided are diagnostic methods for diagnosing and treating, preventing and/or prognosing tissue specific disorders, including cancer, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying agonists and antagonists of cancer antigens of the invention. The present invention further relates to methods and/or compositions for inhibiting the production and/or function of the polypeptides of the present invention.

Full Title	Citation Front	Review Classificatio	n Date Reference	Sequences	Attachments Clair	ns KWWC Drawa Desi
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1... 12. Document ID: US 6727063 B1

L12: Entry 12 of 14

File: USPT Apr 27, 2004

US-PAT-NO: 6727063

DOCUMENT-IDENTIFIER: US 6727063 B1

TITLE: Single nucleotide polymorphisms in genes

DATE-ISSUED: April 27, 2004

h e b b g ee e f e b ef b e

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Lander; Eric S. Cambridge MA Cargill; Michele Gaithersburg MD Ireland; James S. Gaithersburg MD Bolk; Stacev West Roxbury ΜA Daley; George Q. Weston MA McCarthy; Jeanette J. San Diego CA

US-CL-CURRENT: <u>435/6</u>; <u>435/91.1</u>, <u>435/91.2</u>

ABSTRACT:

The invention provides nucleic acid segments of the human genome, particularly nucleic acid segments from a gene, including polymorphic sites. Allele-specific primers and probes hybridizing to regions flanking or containing these sites are also provided. The nucleic acids, primers and probes are used in applications such as phenotype correlations, forensics, paternity testing, medicine and genetic analysis. A role for the thrombospondin gene(s) in vascular disease is also disclosed. Use of single nucleotide polymorphisms in the thrombospondin gene(s) for diagnosis, prediction of clinical course and treatment response, development of therapeutics and development of cell-culture-based and animal models for research and treatment are disclosed.

4 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 8

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13. Document ID: US 6555321 B1

L12: Entry 13 of 14

File: USPT

Apr 29, 2003

US-PAT-NO: 6555321

DOCUMENT-IDENTIFIER: US 6555321 B1

TITLE: Methods for determining cell responses through EphB receptors

DATE-ISSUED: April 29, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Daniel; Thomas O. Nashville TN Stein; Elke San Francisco CA

US-CL-CURRENT: 435/7.1; 435/334, 435/7.2, 435/7.21, 435/7.8

ABSTRACT:

The present invention provides methods for screening an EphB receptor or an EphB receptor-binding ligand for the ability to promote a selected biological activity when in multimeric form. The invention also provides methods for initiating, promoting, directing, or inhibiting biological activities that involve EphB receptors and/or EphB receptor-binding ligands. The invention further provides compositions that can be used in the foregoing methods.

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8 Claims, 9 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

Full Title Citation Front Review Classification Date Reference Claims KodC Draw Des-

14. Document ID: US 6514497 B1

L12: Entry 14 of 14

File: USPT

Feb 4, 2003

US-PAT-NO: 6514497

DOCUMENT-IDENTIFIER: US 6514497 B1

TITLE: Inhibition of LERK-2-mediated cell adhesion

DATE-ISSUED: February 4, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Briskin; Michael J.

Lexington

MΑ

Zou; Lily

Cambridge

MA

US-CL-CURRENT: 424/143.1; 424/130.1, 424/137.1, 424/141.1, 424/152.1, 424/172.1, 530/387.1, 530/387.5, 530/388.1, 530/388.22

ABSTRACT:

Methods of modulating LERK-2-mediated cell adhesion, as well as methods of modulating angiogenesis and inflammation are described. Also described are agents such as antibodies which can modulate LERK-2-mediated cell adhesion, as well as methods of treating angiogenic diseases and inflammatory diseases.

8 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 8

Full Title Citation Front Review	Classification Date Reference Classification Clasms MMC Draw Desi
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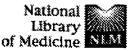
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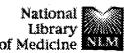
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Physiological, anatomical and genetic identification of CPG neurons in the

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developing mammalian spinal cord.

Prog Neurobiol. 2003 Jul;70(4):347-61. Review.

PMID: 12963092 [PubMed - indexed for MEDLINE]

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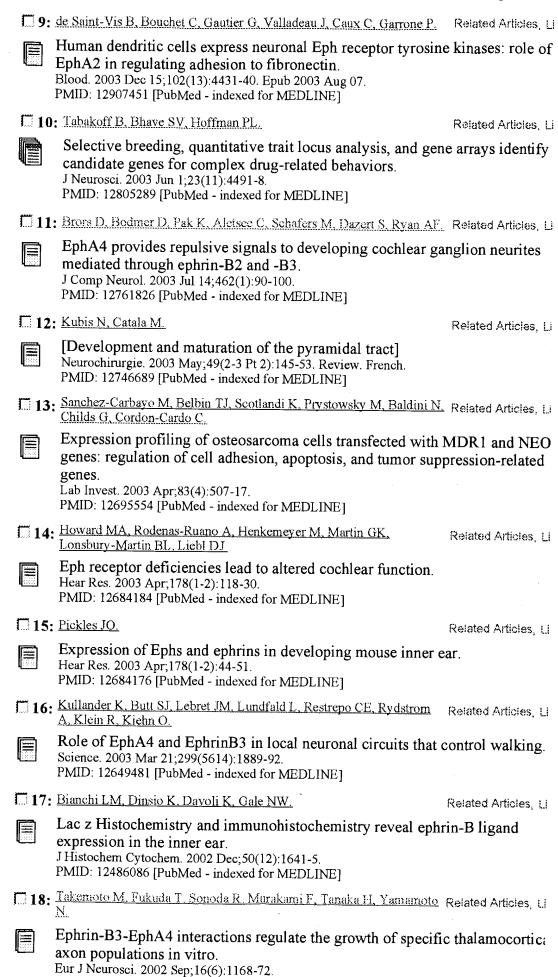
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The receptor tyrosine kinase EphB4 and ephrin-B ligands restrict angiogenic growth of embryonic veins in Xenopus laevis.

Helbling PM, Saulnier DM, Brandli AW.

Institute of Cell Biology, Swiss Federal Institute of Technology, ETH-Honggerberg, CH-8093 Zurich, Switzerland.

The cues and signaling systems that guide the formation of embryonic blood vessels in tissues and organs are poorly understood. Members of the Eph famil of receptor tyrosine kinases and their cell membrane-anchored ligands, the ephrins, have been assigned important roles in the control of cell migration dur embryogenesis, particularly in axon guidance and neural crest migration. Here investigated the role of EphB receptors and their ligands during embryonic blo vessel development in Xenopus laevis. In a survey of tadpole-stage Xenopus embryos for EphB receptor expression, we detected expression of EphB4 receptors in the posterior cardinal veins and their derivatives, the intersomitic veins. Vascular expression of other EphB receptors, including EphB1, EphB2 EphB3, could however not be observed, suggesting that EphB4 is the principal EphB receptor of the early embryonic vasculature of Xenopus. Furthermore, w found that ephrin-B ligands are expressed complementary to EphB4 in the somites adjacent to the migratory pathways taken by intersomitic veins during angiogenic growth. We performed RNA injection experiments to study the function of EphB4 and its ligands in intersomitic vein development. Disruption EphB4 signaling by dominant negative EphB4 receptors or misexpression of ephrin-B ligands in Xenopus embryos resulted in intersomitic veins growing abnormally into the adjacent somitic tissue. Our findings demonstrate that Eph and B-class ephrins act as regulators of angiogenesis possibly by mediating repulsive guidance cues to migrating endothelial cells.

PMID: 10603345 [PubMed - indexed for MEDLINE]

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Role of EphA4 and EphrinB3 in local neuronal circuits that contr walking.

Kullander K, Butt SJ, Lebret JM, Lundfald L, Restrepo CE, Rydstrom A Klein R, Kiehn O.

Department of Medical Biochemistry, Gothenburg University, Medicinaregata A, 405 30 Gothenburg, Sweden. klas.kullander@medkem.gu.se

Local circuits in the spinal cord that generate locomotion are termed central pattern generators (CPGs). These provide coordinated bilateral control over the normal limb alternation that underlies walking. The molecules that organize th mammalian CPG are unknown. Isolated spinal cords from mice lacking either EphA4 receptor or its ligand ephrinB3 have lost left-right limb alternation and instead exhibit synchrony. We identified EphA4-positive neurons as an excitat component of the locomotor CPG. Our study shows that dramatic locomotor changes can occur as a consequence of local genetic rewiring and identifies ge required for the development of normal locomotor behavior.

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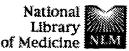
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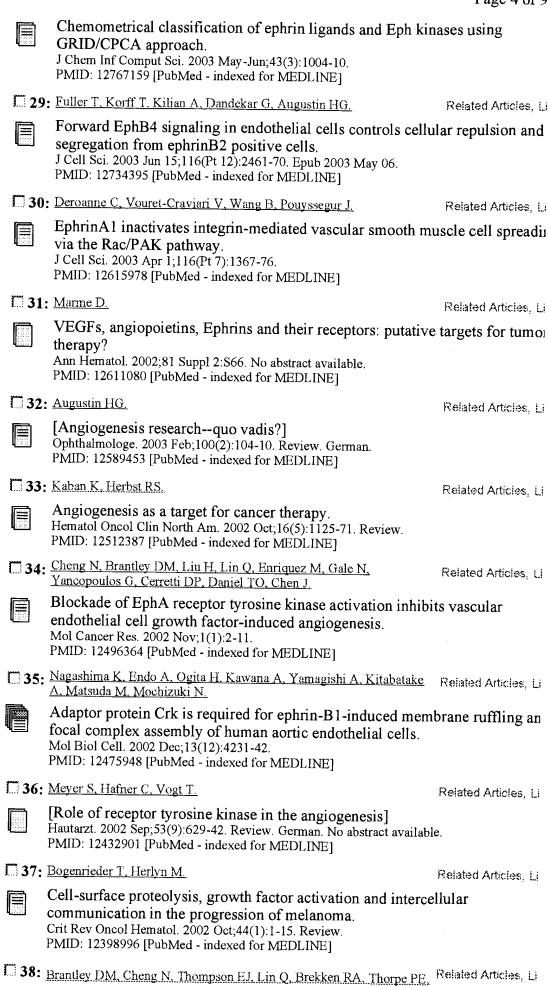
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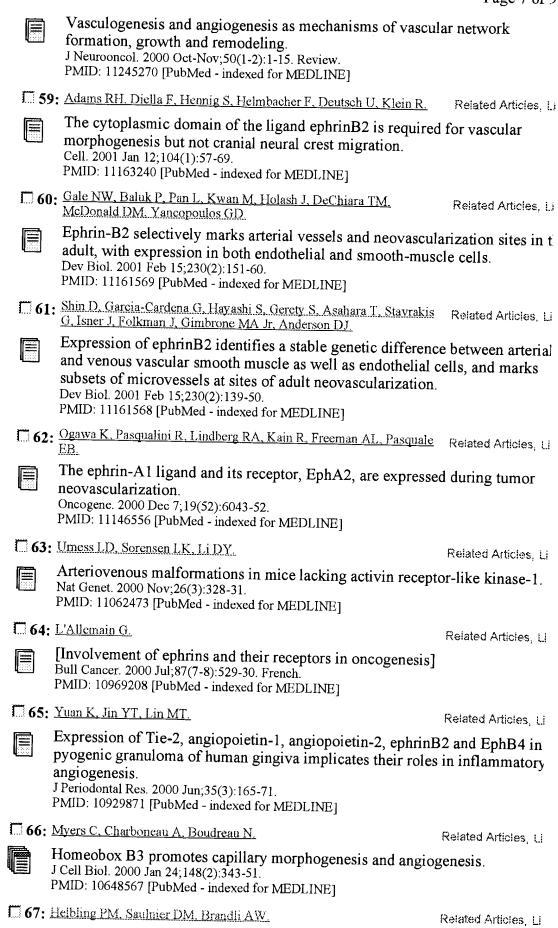
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Roles of ephrinB ligands and EphB receptors in cardiovascular development: demarcation of arterial/venous domains, vascular morphogenesis, and sprouting angiogenesis.

Adams RH, Wilkinson GA, Weiss C, Diella F, Gale NW, Deutsch U, Risau W, Klein R.

European Molecular Biology Laboratory, D-69117 Heidelberg, Germany.

Eph receptor tyrosine kinases and their cell-surface-bound ligands, the ephrins regulate axon guidance and bundling in the developing brain, control cell migration and adhesion, and help patterning the embryo. Here we report that tv ephrinB ligands and three EphB receptors are expressed in and regulate the formation of the vascular network. Mice lacking ephrinB2 and a proportion of double mutants deficient in EphB2 and EphB3 receptor signaling die in utero before embryonic day 11.5 (E11.5) because of defects in the remodeling of the embryonic vascular system. Our phenotypic analysis suggests complex interactions and multiple functions of Eph receptors and ephrins in the embryo vasculature. Interaction between ephrinB2 on arteries and its EphB receptors o veins suggests a role in defining boundaries between arterial and venous doma Expression of ephrinB1 by arterial and venous endothelial cells and EphB3 by veins and some arteries indicates that endothelial cell-to-cell interactions between ephrins and Eph receptors are not restricted to the border between arteries and veins. Furthermore, expression of ephrinB2 and EphB2 in mesenchyme adjace to vessels and vascular defects in ephB2/ephB3 double mutants indicate a requirement for ephrin-Eph signaling between endothelial cells and surroundir. mesenchymal cells. Finally, ephrinB ligands induce capillary sprouting in vitro with a similar efficiency as angiopoietin-1 (Angl) and vascular endothelial growth factor (VEGF), demonstrating a stimulatory role of ephrins in the remodeling of the developing vascular system.

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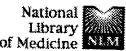
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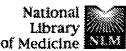
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Vasculogenesis and angiogenesis as mechanisms of vascular network formation, growth and remodeling.

Patan S.

Division of Cardiology, Albert Einstein College of Medicine, Yeshiva Univers Bronx, New York 10461, USA. spatan@aecom.yu.du

Two distinct mechanisms, vasculogenesis and angiogenesis implement the formation of the vascular network in the embryo. Vasculogenesis gives rise to heart and the first primitive vascular plexus inside the embryo and in its surrounding membranes, as the yolk sac circulation. Angiogenesis is responsib for the remodeling and expansion of this network. While vasculogenesis refers in situ differentiation and growth of blood vessels from mesodermal derived hemangioblasts, angiogenesis comprises two different mechanisms: endothelia sprouting and intussusceptive microvascular growth (IMG). The sprouting process is based on endothelial cell migration, proliferation and tube formation IMG divides existing vessel lumens by formation and insertion of tissue folds columns of interstitial tissue into the vessel lumen. The latter are termed interstitial or inter-vascular tissue structures (ITSs) and tissue pillars or posts. Intussusception also includes the establishment of new vessels by in situ loop formation in the wall of large veins. The molecular regulation of these distinct mechanisms is discussed in respect to the most important positive regulators, vascular endothelial growth factor (VEGF) and its receptors flk-1 (KDR) and i 1, the Angiopoietin/tie system and the ephrin-B/EpH-B system. The cellular mechanisms and the molecular regulation of angiogenesis in the pathological state are summarized and the differences of physiological and pathological angiogenesis elaborated.

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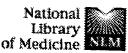
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Eph receptors and ephrin ligands. essential mediators of vascular development.

Adams RH, Klein R.

Developmental Biology Program, Heidelberg, Germany.

The molecular and cellular mechanisms governing vascular development are s poorly understood. Prominent among the intercellular signals that control the initial establishment of the vascular network (termed vasculogenesis) and the subsequent remodeling process (called angiogenesis) are soluble ligands that signal through receptor tyrosine kinases (RTKs). Recent reports have added ce bound ephrin ligands and their cognate Eph RTKs to the list of key players in vascular development.

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Angiogenesis factors.

Kuwano M, Fukushi J, Okamoto M, Nishie A, Goto H, Ishibashi T, Ono N

Department of Medical Biochemistry, Graduate School of Medical Sciences, Kyushu University, Fukuoka.

Angiogenesis is a recent highlight in the medical field; the developmental proc and pathological conditions for various diseases can be understood from the no aspect of "angiogenesis". Many angiogenesis-related factors are involved in th development of vessels during embryogenesis (vasculogenesis), as well as the induction of new vessels in response to physiological or pathological stimuli. I particular, the appearance of hemangioblasts, precursor cells for vascular endothelial cells and blood cells, and blood islands are expected to play a "prelude" role in tubulogenesis. Gene knock out mice of vascular endothelial growth factor (VEGF)/VEGF receptor, ephrin-B2, and angiopoietin-1 results is failure of normal vessels production. Dormant factors derived from proteolytic cleavage of various physiological substrates are expected to balance a homeostasis of "angiogenic states" in the host. Furthermore, angiogenesis unde various pathological conditions of malignant tumors, ocular diseases, psoriasis rheumatoid arthritis, atherosclerosis and other diseases is associated with comp angiogenesis networks, suggesting pleiotropic mechanisms for angiogenesis.

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The ephrins and Eph receptors in angiogenesis.

Cheng N, Brantley DM, Chen J.

Department of Cancer Biology, Vanderbilt University School of Medicine, A-4323 MCN, 1161 21st Avenue South, Nashville, TN 37232, USA.

Eph receptors are a unique family of receptor tyrosine kinases that play critical roles in embryonic patterning, neuronal targeting, vascular development and at neovascularization. Engagement of Eph receptors by ephrin ligands mediates critical steps of angiogenesis, including juxtacrine cell-cell contacts, cell adhes to extracellular matrix, and cell migration. Recent evidence from in vitro angiogenesis assays and analysis of mice deficient for one or more members of the Eph family establishes the role of Eph signaling in sprouting angiogenesis blood vessel remodeling during vascular development. Furthermore, elevated expression of Eph receptors and ephrin ligands is associated with tumors and associated tumor vasculature, suggesting that Eph receptors and their ephrin ligands also play critical roles in tumor angiogenesis and tumor growth. This review will focus on the relevance of Eph receptor signaling in embryonic and adult neovascularization, and possible contributions to tumor growth and metastasis.

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Augustin HG.

Abt. fur Vasklare Biologie und Angiogeneseforshcung, Klinik fur Tumorbiolo Freiburg. augustin@angiogenese.de

The field of angiogenesis research has seen an explosion of knowledge within last 10 years. More than 3500 angiogenesis-related papers are presently being published per year compared to the less than 200 annual papers published in the early 1990s. Paralleling the progress in the field of basic angiogenesis research, translational research has led to the identification of more than 100 angiomanipulatory compounds. Presently, more than 40 substances are in variable phases of clinical trials. The prospect of these exciting developments is present dampened by the negative outcome of some advanced clinical trials. Thus, following euphoria and disillusion, the field is presently experiencing that translational clinical research requires endurance to eventually accomplish the successful implementation of angiomanipulatory therapies in the clinical settin. The present article provides an overview of the field of angiogenesis research a summarizes ongoing efforts aimed at developing angiomanipulatory therapies.

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Elk-L3, a novel transmembrane ligand for the Eph family of receptor tyrosine kinases, expressed in embryonic floor plate, roo plate and hindbrain segments.

Gale NW, Flenniken A, Compton DC, Jenkins N, Copeland NG, Gilbert D Davis S, Wilkinson DG, Yancopoulos GD.

Regeneron Pharmaceuticals, Inc, Tarrytown, New York 10591, USA.

The Eph family of receptor tyrosine kinases has 13 distinct members and seven ligands for these receptors have been described to date. These receptors and th ligands have been implicated in regulating neuronal axon guidance and in patterning of the developing nervous system and may also serve a patterning a compartmentalization role outside of the nervous system as well. The ligands a all membrane-attached, and this attachment appears to be crucial for their norn function; five of the known ligands are linked to the membrane via a glycosyl phosphotidylinositol (GPI) linkage, while two of the ligands are transmembran proteins. Despite the large number of Eph family receptors and ligands, they ca be divided into just two major subclasses based on their binding specificities. the GPI-anchored ligands bind and activate one subclass of the Eph receptors (that represented by Eck) while the two transmembrane ligands bind and active the other major subclass of receptors (represented by Elk). Here we report the identification and characterization of the third, and most divergent, member of transmembrane group of Eph ligands, which we term Elk-L3 (Elk-related received ligand number 3). Elk-L3 is notable for its remarkably restricted and prominen expression in the floor plate and roof plate of the developing neural tube and it rhombomere-specific expression in the developing hindbrain. The Elk-L3 gene has been localized to mouse chromosome 11 and human chromosome 17.

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Ephrin-B3, a ligand for the receptor EphB3, expressed at the midline of the developing neural tube.

Bergemann AD, Zhang L, Chiang MK, Brambilla R, Klein R, Flanagan J

Department of Cell Biology, Harvard Medical School, Boston, Massachusetts 02115, USA.

The ephrins are a family of ligands that bind to Eph family receptor tyrosine kinases, and have been implicated in axon guidance and other patterning processes during vertebrate development. We describe here the identification 2 characterization of murine ephrin-B3. The cDNA encodes a 340 amino acid transmembrane molecule, most closely related to the two other known transmembrane ligands, ephrin-B1 and ephrin-B2. In addition to homology in their extracellular receptor binding domains, these transmembrane ligands shall striking homology between their cytoplasmic domains, with 31 of the last 34 amino acids of ephrin-B3 being identical to ephrin-B2, suggesting functional interactions of the cytoplasmic tail. While most Eph family ligands are promiscuous in their interactions with Eph receptors, binding studies with the receptors known to bind other transmembrane ligands only revealed a high affinity interaction of ephrin-B3 with EphB3, with a dissociation constant of approximately 1 nM. In situ hybridization of mouse embryos showed ephrin-B is expressed prominently at the dorsal and ventral midline of the neural tube, particularly in the floor plate, a structure with key functions in patterning the nervous system. The isolation of this ligand may help to elucidate the molecula basis of patterning activities at the neural tube midline.

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Graduate School of Medicine, Osaka University, Suita, Osaka, Japan Hepatology, (October 2003) Vol. 38, No. 4 Suppl. 1, pp. 760A-761A. print. Meeting Info.: 54th Annual Meeting of the American Association for the Study of Liver Diseases. Boston, MA, USA. October 24-28, 2003. American Association for the Study of Liver Diseases.

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ANSWER 3 OF 28 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on L4AN 2003:543156 BIOSIS DN PREV200300538665 EPHRINA1 SIGNALING INHIBITS VEGF - INDUCED ERK1/2 PHOSPHORYLATION AND TΙ RETINAL ENDOTHELIAL CELL PROLIFERATION. Ojima, T. [Reprint Author]; Takagi, H. [Reprint Author]; Suzuma, K. [Reprint Author]; Oh, H. [Reprint Author]; Suzuma, I. [Reprint Author]; Ohashi, H. [Reprint Author]; Watanabe, D. [Reprint Author]; Suganami, E. [Reprint Author]; Honda, Y. [Reprint Author]
Ophthalmology and Visual Sciences, Kyoto University Graduate School of ΑU CS Medicine, Kyoto, Japan ARVO Annual Meeting Abstract Search and Program Planner, (2003) Vol. 2003, SO pp. Abstract No. 2881. cd-rom. Meeting Info.: Annual Meeting of the Association for Research in Vision and Ophthalmology. Fort Lauderdale, FL, USA. May 04-08, 2003. Association for Research in Vision and Ophthalmology. Conference; (Meeting) Conference; Abstract; (Meeting Abstract) DT LΑ English Entered STN: 19 Nov 2003 ED Last Updated on STN: 19 Nov 2003 ANSWER 4 OF 28 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on L4STNAN 2003:542402 BIOSIS ***Ephrin*** B1 is expressed on human luteinizing granulosa cells in corpora lutea of the early luteal phase: The possible involvement of the B class Eph- ***ephrin*** system during corpus luteum formation. Egawa, Miho; Yoshioka, Shinya; Higuchi, Toshihiro; Sato, Yukiyasu; Tatsumi, Keiji; Fujiwara, Hiroshi [Reprint Author]; Fujii, Shingo Department of Gynecology and Obstetrics, Faculty of Medicine, Kyoto University, Sakyo-ku, Kyoto, 606-8507, Japan fuji@kuhp.kyoto-u.ac.jp Journal of Clinical Endocrinology & Metabolism, (September 2003) Vol. 88, No. 9, pp. 4384-4392. print. ISSN: 0021-972X (ISSN print). Article PREV200300543738 DN TIΑU CS SO DT Article English LΑ Entered STN: 19 Nov 2003 EDLast Updated on STN: 19 Nov 2003 ANSWER 5 OF 28 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on L4STN 2002:214449 BIOSIS AN PREV200200214449 DNCoexpression of ***ephrin*** -Bs and their receptors in colon ΤI carcinoma. Liu, Wenbiao; Ahmad, Syed A.; Jung, Young D.; Reinmuth, Niels; Fan, Fan; Bucana, Corazon D.; Ellis, Lee M. [Reprint author] AU Department of Surgical Oncology, University of Texas M. D. Anderson Cancer Center, 1515 Holcombe Boulevard, 444, Houston, TX, 77030-4009, USA CS lellis@mdanderson.org Cancer, (February 15, 2002) Vol. 94, No. 4, pp. 934-939. print. CODEN: CANCAR. ISSN: 0008-543X. SO DT Article LА English Entered STN: 27 Mar 2002 ED Last Updated on STN: 27 Mar 2002 CAPLUS COPYRIGHT 2004 ACS on STN L4ANSWER 6 OF 28 2004:355085 ΑN CAPLUS DN140:369944 Human tissue-specific housekeeping genes identified by expression TI profiling Aburatani, Hiroyuki; Yamamoto, Shogo INNGK Insulators, Ltd., Japan PCT Int. Appl., 372 pp. PA SO CODEN: PIXXD2 DT Patent

APPLICATION NO.

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Japanese

PATENT NO.

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     Alitalo, Kari; Kubo, Hajime
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     Comparative analysis of embryonic gene expression defines potential
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     Helbling, Paul M.; Saulnier, Didier M. E.; Robinson, Vicky; Christiansen, Jeff H.; Wilkinson, David G.; Brandli, Andre W.
AU
     Institute of Cell Biology, Swiss Federal Institute of Technology, Zurich,
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     CH-8093, Switz.
     Developmental Dynamics (1999),
                                         216(4/5), 361-373
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        ***Ephrins***
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     Blits-Huizinga C.T.; Nelersa C.M.; Malhotra A.; Liebl D.J. D.J. Liebl, Miami Project to Cure Paralysis, Univ. of Miami School of
AU
CS
     Medicine, 1095 NW 14th Terrace, Miami, FL 33136, United States.
     dliebl@miami.edu
SO
      IUBMB Life, (2004) 56/5 (257-265).
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      Pasquale E.B.
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      E.B. Pasquale, Burnham Institute, San Diego, CA 92037, United States.
CS
      elenap@burnham.org
      Nature Neuroscience, (2004) 7/5 (417-418).
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      Functions of EPH Receptors and
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      Principal Investigator: IKEGAKI, NAOHĪKO; IKEGAKI@EMAIL.CHOP.EDU,
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      Prognostic significance of EPHB6, EFNB2 and EFNB3 expressions in
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      Tang X X; Zhao H Q; Robinson M E; Cnaan A; London W; Cohn S L; Cheung N K V; Brodeur G M; Evans A E; Ikegaki N (Reprint)
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      NY 10021
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SO
      NY 10158-0012.
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         Novel agents that modulate Eph receptor activity Pasquale, Elena B., San Diego, CA, UNITED STATES Koolpe, Mitchell, San Diego, CA, UNITED STATES Murai, Keith K., Candiac, CANADA
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        Compositions and methods for regulating the kinase domain of receptor
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        Sicheri, Frank, Toronto, CANADA
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Aziz, Natasha, Palo Alto, CA, UNITED STATES
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        ICS: C07H021-04; C12P021-02; C12N005-06; C07K014-47
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 23 OF 28 USPATFULL on STN
L4
        2004:2047
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AN
TI
        Breast cancer progression signatures
        Erlander, Mark G., Encinitas, CA, UNITED STATES
IN
        Ma, Xia-Jun, San Diego, CA, UNITED STATES
        Sqroi, Dennis C., Winchester, MA, UNITED STATES
                                   20040101
ΡI
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                             A1
                                   20011221 (10)
ΑI
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INCL
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               435/287.200; 702/020.000
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        ICM: C120001-68
        ICS: G06F019-00; G01N033-48; G01N033-50; C12M001-34
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 24 OF 28
L4
                        USPATFULL on STN
        2003:220740
AN
                      USPATFULL
        Methods and compositions for diagnosing and treating rheumatoid
ΤI
        arthritis
        Pittman, Debra D., Windham, NH, UNITED STATES
ΙN
        Feldman, Jeffrey L., Arlington, MA, UNITED STATES
        Shields, Kathleen M., Harvard, MA, UNITED STATES
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20030814
PΙ
        US 2003154032
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 25 OF 28 USPATFULL on STN
L4
ΑN
        2003:120026 USPATFULL
        Identification of modulatory molecules using inducible promoters Brown, Steven J., San Diego, CA, UNITED STATES
TI
IN
                                                CA, UNITED STATES
        Dunnington, Damien J., San Diego,
                Imran, San Diego, CA, UNITED STATES 8082511 A1 20030501
        Clark,
PI
        US 2003082511
                                    20010925
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        US 2001-965201
                              Α1
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APPLICATION
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 26 OF 28
                         USPATFULL on STN
L4
        2003:30255 USPATFULL
AN
                              regulation of G-protein coupled chemoattraction,
TI
            ***ephrin***
        compositions, and methods of use
        Flanagan, John G., Newton, MA, UNITED STATES
Lu, Qiang, Brookline, MA, UNITED STATES
Sun, Edna E., Brookline, MA, UNITED STATES
US 2003022202 A1 20030130
IN
        US 2003022202
PΙ
AΙ
        US 2002-113794
                              Α1
                                     20020401
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        US 2001-280260P
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        2002:266423
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AN
TI
        Peptides that modulate the interaction of B class
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                                                                                         and
        PDZ domains
IN
        Lin, Danny,
                      Scarborough, CANADA
        Pawson, Anthony, Toronto, CANADA
                                     CANADA
               Gerald, East York,
        US 2002147306
                                     20021010
PΙ
                              A1
                              A1
                                     20010521 (9)
        US 2001-862179
ΑI
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        WO 1999-CA1101
        US 1998-109158P
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ICM: C07K014-435
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 28 OF 28 USPATFULL on STN
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       2002:99407 USPATFULL
AN
       Nucleic acids, proteins and antibodies
TI
       Rosen, Craig A., Laytonsville, MD, UNITED STATES
IN
               Steven M., Olney, MD, UNITED STATES
052308 A1 20020502
ΡI
       US 2002052308
       US 2001-925301
                            A1
                                  20010810 (9)
ΑI
RLI
       Continuation of Ser. No. WO 2000-US5882, filed on 8 Mar 2000, UNKNOWN
       US 1999-124270P
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NCL
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       NCLS:
               530/350.000; 435/320.100; 435/325.000
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        ICM: A61K031-00
        ICS: C12Q001-68; G01N033-53; C07H021-04; C12N009-00; C07K014-435;
        C12N005-06
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
=> S ephrin-B3 OR ephrinB3
  51 FILES SEARCHED...
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DUPLICATE IS NOT AVAILABLE IN 'ADISINSIGHT, ADISNEWS, BIOCOMMERCE, DGENE,
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MEDICONF, NUTRACEUT, PCTGEN, PHAR, PHARMAML, PROUSDDR, RDISCLOSURE, SYNTHLINE'. ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE
PROCESSING COMPLETED FOR L6
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L7
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     ANSWER 1 OF 159
                        CAPLUS
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L7
     2004:355085
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AN
DN
     140:369944
     Human tissue-specific housekeeping genes identified by expression
TI
     profiling
IN
     Aburatani, Hiroyuki; Yamamoto, Shogo
     NGK Insulators, Ltd., Japan PCT Int. Appl., 372 pp.
PA
SO
     CODEN: PIXXD2
DT
     Patent
     Japanese
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     PATENT NO.
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     WO 2004035785
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                THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
                        CAPLUS
                                 COPYRIGHT 2004 ACS on STN
     ANSWER 2 OF 159
L7
     2004:204003
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AN
      140:248255
DN
     Synovial sarcoma up-regulated gene FZD10 (Frizzled homolog 10) and other
ΤI
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Nakamura, Yusuke; Katagiri, Toyomasa
IN
      Oncotherapy Science, Inc., Japan; Japan as Represented by President of the
PA
      University of Tokyo
      PCT Int. Appl., 143 pp.
SO
      CODEN: PIXXD2
DT
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LA English FAN.CNT 1
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      PATENT NO.
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PRAI US 2002-407506P
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                                     20030711
      US 2003-486195P
      ANSWER 3 OF 159 USPATFULL on STN
L7
        2004:233749 USPATFULL
AN
        Novel agents that modulate Eph receptor activity Pasquale, Elena B., San Diego, CA, UNITED STATES Koolpe, Mitchell, San Diego, CA, UNITED STATES
TI
IN
        Murai, Keith K., Candiac, CANADA
                                    20040916
PI
        US 2004180823
                             A1
                                               (10)
ΑI
        US 2003-652407
                             Α1
                                    20030829
        US 2002-413242P
                              20020924 (60)
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        ICM: A61K038-17
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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        2004:233299 USPATFULL
NA
TI
        Genetic diagnosis of alcoholism subtypes
        Tabakoff, Boris, Denver, CO, UNITED STATES
IN
        Martinez, Larry, Colorado Springs, CO, UNITED STATES
Hoffman, Paula, Denver, CO, UNITED STATES
THE REGENTS OF THE UNIVERSITY OF COLORADO, a body corporate, Boulder, CA
PA
        (U.S. corporation)
        US 2004180370
US 2004-766590
US 2003-443072F
ΡI
                              A1
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AΙ
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            2003-443072P
PRAI
                              20030127 (60)
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        ICM: C12Q001-68
        ICS: C12P019-34
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
      ANSWER 5 OF 159 USPATFULL on STN
        2004:196424 USPATFULL
AN
        Lectin compositions and methods for modulating an immune response to an
TI
        antigen
        Segal, Andrew H., Boston, MA, UNITED STATES
IN
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PA
       Genitrix, LLC (U.S. corporation)
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       US 2004151728
PI
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AI
                                  20030919
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       US 2003-666834
       Division of Ser. No. US 2003-645000, filed on 20 Aug 2003, PENDING
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       US 2002-404823P
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PRAI
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APPLICATION
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 6 OF 159 USPATFULL on STN
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                     USPATFULL
AN
       2004:177819
TI
       Methods for inhibiting angiogenesis by EphB receptor antagonists
       Aguet, Michel, Lutry, SWITZERLAND
IN
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PI
       US 2004136983
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AI
       US 2004-770543
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                                  20040202
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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     ANSWER 7 OF 159
                       USPATFULL on STN
       2004:172470 USPATFULL
AN
TI
       Compositions and methods for regulating the kinase domain of receptor
       tyrosine kinases
       Sicheri, Frank, Toronto, CANADA
Wybenga-Groot, Leanne, Etobicoke, CANADA
Pawson, Tony, Toronto, CANADA
IN
       US 2004132634
PΙ
                            A1
                                  20040708
                            Α1
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AI
       US 2004-470840
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       WO 2002-CA114
                                  20020131
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       US 2001-60265510
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       ICM: A61K031-00
       ICS: G06F019-00; G01N033-48; G01N033-50; C12N009-12
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 8 OF 159
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       2004:165307
                     USPATFULL
AN
ΤI
       Lectin compositions and methods for modulating an immune response to an
       antigen
IN
       Segal, Andrew H., Boston, MA, UNITED STATES
       Young, Elihu, Sharon, MA, UNITED STATES
       Genitrix, LLC (U.S. corporation)
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       US 2004126793
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m PI}
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ΑI
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PRAI
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         NCLM:
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         NCLS:
                  530/395.000; 536/023.500
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         ICM: C12Q001-68
         ICS: C07H021-04; C07K014-47; C07K014-415; C12N005-04
     INDEXING IS AVAILABLE FOR THIS PATENT.
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      ANSWER 9 OF 159 USPATFULL on STN
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         2004:164872 USPATFULL
AN
         Lectin compositions and methods for modulating an immune response to an
TI
         antigen
         Segal, Andrew H., Boston, MA, UNITED STATES
IN
         Young, Elihu, Sharon, MA, UNITED STATES Genitrix, LLC (U.S. corporation)
PΑ
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         US 2004126357
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         Division of Ser. No. US 2003-645000, filed on 20 Aug 2003, PENDING US 2002-404823P 20020820 (60)
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US 2003-487407P
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         ICS: A61K039-00; A61K038-19
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 10 OF 159
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                              USPATFULL on STN
         2004:133338 USPATFULL
AN
         Targets for therapeutic intervention identified in the mitochondrial
TI
         proteome
         Ghosh, Soumitra S., San Diego, CA, UNITED STATES Fahy, Eoin D., San Diego, CA, UNITED STATES Zhang, Bing, Spring, TX, UNITED STATES Gibson, Bradford W., Berkeley, CA, UNITED STATES Taylor, Steven W., San Diego, CA, UNITED STATES Glenn, Gary M., Encinitas, CA, UNITED STATES Warnock Dale E. San Diego, CA UNITED STATES
IN
         Warnock, Dale E., San Diego, CA, UNITED STATES
Gaucher, Sara P., Castro Valley, CA, UNITED STATES
MitoKor Inc., San Diego, CA, UNITED STATES, 92121 (U.S. corporation)
PA
         The Buck Institute for Age Research, Novato, CA, UNITED STATES,
         94948-0638 (U.S. corporation)
         US 2004101874
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 11 OF 159 USPATFULL on STN
L7
         2004:101093 USPATFULL
AN
         Methods of diagnosis of bladder cancer, compositions and methods of
TI
         screening for modulators of bladder cancer
         Mack, David H., Menlo Park, CA, UNITED STATES
Aziz, Natasha, Palo Alto, CA, UNITED STATES
Eos Biotechnology, Inc., South San Francisco, CA, UNITED STATES,
IN
PA
         94080-7019 (U.S. corporation)
         US 2004076955
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PΙ
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         US 2002-188832
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                                  20020412 (60)
PRAI
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        US 2001-310099P
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        US 2001-302814P
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        ICM: C12Q001-68
        ICS: C07H021-04; C12P021-02; C12N005-06; C07K014-47
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 12 OF 159
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L7
        2004:76614
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ΑN
        Use of novel stem cell markers for isolation of intestinal stem cells,
TI
        and use of the intestinal stem cells thus obtained for the preparation
        of a therapeutical composition
Clevers, Johannes Carolus, Huis Ter Heide, NETHERLANDS
IN
        Gomez, Eduard Batlle, Utrecht, NETHERLANDS
        Van De Wetering, Marcus Lambertus, Houten, NETHERLANDS
        Suils, Elena Sancho, Utrecht, NETHERLANDS
        Kylix B.V., Driebergen, NETHERLANDS, NL-3971 JD (non-U.S. corporation)
PA
        US 2004058392
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ΑI
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      ANSWER 13 OF 159
L7
        2004:2047 USPATFULL
AN
ΤI
        Breast cancer progression signatures
        Erlander, Mark G., Encinitas, CA, UNITED STATES
IN
        Ma, Xia-Jun, San Diego, CA, UNITED STATES
        Sgroi, Dennis C., Winchester, MA, UNITED STATES US 2004002067 A1 20040101
        UŠ 2004002067
ΡI
        US 2001-28018
                                    20011221 (10)
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                              A1
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APPLICATION
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IC
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ICS: G06F019-00; G01N033-48; G01N033-50; C12M001-34 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 14 OF 159 USPATFULL on STN 2004:103677 USPATFULL
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AN
TI
        Single nucleotide polymorphisms in genes
                Eric S., Cambridge, MA, United States
IN
        Lander
        Cargill, Michele, Gaithersburg, MD, United States
        Ireland, James S., Gaithersburg, MD, United States
Bolk, Stacey, West Roxbury, MA, United States
Daley, George Q., Weston, MA, United States
        McCarthy, Jeanette J., San Diego, CA, United States
Millennium Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
PA
        corporation)
        Whitehead Institute for Biomedical Research, Cambridge, MA, United
        States (U.S. corporation)
        US 6727063
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NCLM: 435/006.000
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IC
          ICM: C12Q001-68
          ICS: C12P019-34
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
                                EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS
       ANSWER 15 OF 159
L7
       RESERVED. on STN
AN
       2004382596
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      Favorable neuroblastoma genes and molecular therapeutics of neuroblastoma. Tang X.X.; Robinson M.E.; Riceberg J.S.; Kim D.Y.; Kung B.; Titus T.B.; Hayashi S.; Flake A.W.; Carpentieri D.; Ikegaki N. N. Ikegaki, Division of Hematology/Oncology, Department of Pediatrics, Emory University School of Medicine, 2040 Ridgewood Drive NE, Atlanta, GA 30322 United States neo ikegaki@07. red emory odu
TΙ
ΑU
CS
       30322, United States. nao_ikegaki@oz.ped.emory.edu
Clinical Cancer Research, (1 Sep 2004) 10/17 (5837-5844).
SO
       Refs: 16
       ISSN: 1078-0432 CODEN: CCREF4
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       ANSWER 16 OF 159 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation.
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       2004:521594
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AN
       The Genuine Article (R) Number: 823ZZ
GΑ
       Rescuing transient corticospinal terminations and promoting growth with
TI
       corticospinal stimulation in kittens
AU
       Salimi I; Martin J H (Reprint)
       Columbia Univ, Ctr Neurobiol & Behav, New York State Psychiat Inst, 1051 Riverside Dr, New York, NY 10032 USA (Reprint); Columbia Univ, Ctr
CS
       Neurobiol & Behav, New York State Psychiat Inst, New York, NY 10032 USA
CYA
       USA
       JOURNAL OF NEUROSCIENCE, (26 MAY 2004) Vol. 24, No. 21, pp. 4952-4961. Publisher: SOC NEUROSCIENCE, 11 DUPONT CIRCLE, NW, STE 500, WASHINGTON, DC
SO
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ISSN: 0270-6474.
DT
       Article; Journal
LΑ
       English
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       *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*
       ANSWER 17 OF 159
                                EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS
L7
       RESERVED. on STN
       2004413235
AΝ
                       EMBASE
      Identification of tissue-restricted transcripts in human islets.
Maffei A.; Liu Z.; Witkowski P.; Moschella F.; Del Pozzo G.; Liu E.;
Herold K.; Winchester R.J.; Hardy M.A.; Harris P.E.
Dr. P.E. Harris, Department of Medicine BB 20-06, Columbia University,
TI
AU
CS
       College of Physicians and Surgeons, 650 West 168th Street, New York, NY
       10032, Italy. pehl@columbia.edu
       Endocrinology, (2004) 145/10 (4513-4521).
SO
       Refs: 64
       ISSN: 0013-7227 CODEN: ENDOAO
CY
       United States
       Journal; Article
DT
FS
                  Endocrinology
       003
                  Human Genetics
       022
                  Clinical Biochemistry
       029
       English
LΑ
       English
\operatorname{SL}
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BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. ANSWER 18 OF 159 1.7 STN DUPLICATE 1 AN2004:192651 BIOSIS DNPREV200400180545 ΤI Abnormal hippocampal axon bundling in EphB receptor mutant mice. Chen, Zhi-Yong; Sun, Chunhua; Reuhl, Kenneth; Bergemann, Andrew; ΑU Henkemeyer, Mark; Zhou, Renping [Reprint Author]
Department of Chemical Biology, College of Pharmacy, Rutgers University, CS Piscataway, NJ, 08854, USA rzhou@rci.rutgers.edu Journal of Neuroscience, (March 10 2004) Vol. 24, No. 10, pp. 2366-2374. SO print ISSN: 0270-6474 (ISSN print). DT Article English LΑ Entered STN: 7 Apr 2004 ED Last Updated on STN: 7 Apr 2004 MEDLINE on STN L7 ANSWER 19 OF 159 2004233594 MEDLINE ANPubMed ID: 15124102 DN Mutations of the ephrin-B1 gene cause craniofrontonasal syndrome. ΤI Wieland Ilse; Jakubiczka Sibylle; Muschke Petra; Cohen Monika; Thiele ΑU Hannelore; Gerlach Klaus L; Adams Ralf H; Wieacker Peter Institut fur Humangenetik, Otto-von-Guericke-Universitat Magdeburg, 39120 CS Magdeburg, Germany American journal of human genetics, (2004 Jun) 74 (6) 1209-15. SO Journal code: 0370475. ISSN: 0002-9297. CY United States Journal; Article; (JOURNAL ARTICLE) DT LΑ English Priority Journals GENBANK-XM038809 FS OS ΕM 200407 Entered STN: 20040511 ED Last Updated on STN: 20040707 Entered Medline: 20040706 ANSWER 20 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. L7 STN AN2004:316085 BIOSIS PREV200400316690 DNInvasiveness of breast carcinoma cells and transcript profile: Eph TIreceptors and ephrin ligands as molecular markers of potential diagnostic and prognostic application. Fox, Brian P.; Kandpal, Raj P. [Reprint Author] Dept Biol Sci, Fordham Univ, Bronx, NY, 10458, USA AU CS kandpal@fordham.edu Biochemical and Biophysical Research Communications, (June 11 2004) Vol. 318, No. 4, pp. 882-892. print. CODEN: BBRCA9. ISSN: 0006-291X. SO DT Article LA English Entered STN: 15 Jul 2004 ED Last Updated on STN: 15 Jul 2004 COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS L7 ANSWER 21 OF 159 EMBASE RESERVED. on STN 2004385354 AΝ EMBASE Ephrin signaling in axon guidance. TIΑU Huot J. CS Canada. Jacques.Huot@phc.ulaval.ca Progress in Neuro-Psychopharmacology and Biological Psychiatry, (2004) SO 28/5 (813-818). Refs: 24 ISSN: 0278-5846 CODEN: PNPPD7 PUI S 0278-5846(04)00082-X CY United States DTJournal; General Review FS 008 Neurology and Neurosurgery Clinical Biochemistry 029 LА English English SLANSWER 22 OF 159 CAPLUS COPYRIGHT 2004 ACS on STN L7

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DN
      140:301431
      Differential gene expression of Eph receptors and ephrins in benign human
TI
      tissues and cancers
      Hafner, Christian; Schmitz, Gerd; Meyer, Stefanie; Bataille, Frauke; Hau,
AU
      Peter; Langmann, Thomas; Dietmaier, Wolfgang; Landthaler, Michael; Vogt,
      Thomas
      Department of Dermatology, University of Regensburg, Regensburg, Germany Clinical Chemistry (Washington, DC, United States) (2004), 50(3), 490-499
CS
SO
      CODEN: CLCHAU; ISSN: 0009-9147
      American Association for Clinical Chemistry
PB
DT
      Journal
      English
LΑ
RE.CNT
         40
                 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD
                 ALL CITATIONS AVAILABLE IN THE RE FORMAT
                           EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS
      ANSWER 23 OF 159
L7
      RESERVED. on STN
      2004190550
AN
                    EMBASE
      Eph-ephrin promiscuity is now crystal clear. Pasquale E.B.
TI
ΑU
      E.B. Pasquale, Burnham Institute, San Diego, CA 92037, United States.
CS
      elenap@burnham.org
      Nature Neuroscience, (2004) 7/5 (417-418).
SO
      Refs: 15
      ISSN: 1097-6256 CODEN: NANEFN
CY
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DT
      Journal; (Short Survey)
               Neurology and Neurosurgery
Clinical Biochemistry
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      English
LΑ
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L7
      ANSWER 24 OF 159
                           EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS
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      2004407732
AN
                    EMBASE
ΤI
      Ephrins and their receptors: Binding versus biology.
     Blits-Huizinga C.T.; Nelersa C.M.; Malhotra A.; Liebl D.J. D.J. Liebl, Miami Project to Cure Paralysis, Univ. of Miami School of Medicine, 1095 NW 14th Terrace, Miami, FL 33136, United States.
ΑU
CS
      dliebl@miami.edu
      IUBMB Life, (2004) 56/5 (257-265).
SO
      Refs: 49
      ISSN: 1521-6543 CODEN: IULIF8
CY
      United States
DT
      Journal; General Review
FS
      800
               Neurology and Neurosurgery
               Developmental Biology and Teratology
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LΑ
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\operatorname{SL}
L7
      ANSWER 25 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation.
                                                                                            on
                                                                    DUPLICATE 2
      STN
AN
      2004:321300 BIOSIS
DN
      PREV200400319321
      Intramembrane cleavage of ***ephrinB3***
                                                            by the human rhomboid family
TI
      protease, RHBDL2.
      Pascall, John C.; Brown, Kenneth D. [Reprint Author]
Signalling Programme, Babraham Inst, Babraham Hall, Cambridge, CB2 4AT,
ΑU
CS
      England
      ken.brown@bbsrc.ac.uk
SO
      Biochemical and Biophysical Research Communications, (April 23 2004) Vol.
      317, No. 1, pp. 244-252. print. CODEN: BBRCA9. ISSN: 0006-291X.
DT
      Article
LA
      English
      Entered STN: 21 Jul 2004
ED
      Last Updated on STN: 21 Jul 2004
      ANSWER 26 OF 159
                           EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS
L7
      RESERVED. on STN
      2004065640
AN
                    EMBASE
      Ganglion cell axon pathfinding in the retina and optic nerve. Oster S.F.; Deiner M.; Birgbauer E.; Sretavan D.W.
TI
ΑU
      D.W. Sretavan, Depts. of Ophthalmol. and Physiology, Programs Neurosci.
CS
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Francisco, CA 94143, United States. dws@itsa.ucsf.edu Seminars in Cell and Developmental Biology, (2004) 15/1 (125-136). SO Refs: 82 ISSN: 1084-9521 CODEN: SCDBFX CY United Kingdom DTJournal; General Review Neurology and Neurosurgery Ophthalmology Developmental Biology and Teratology FS 012 021 029 Clinical Biochemistry LAEnglish SLEnglish EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS ANSWER 27 OF 159 L7 RESERVED. on STN 2003509599 ΑN **EMBASE** EphB4 signaling is capable of mediating ephrinB2-induced inhibition of TI cell migration. Sturz A.; Bader B.; Thierauch K.-H.; Glienke J. ΑU J. Glienke, Research Laboratories of Schering AG, Berlin, Germany. CS jens.glienke@schering.de Biochemical and Biophysical Research Communications, (2 Jan 2004) 313/1 SO (80-88). Refs: 21 ISSN: 0006-291X CODEN: BBRCA CY United States DT Journal; Article FS Clinical Biochemistry 029 LΑ English SL English ANSWER 28 OF 159 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 3 L7 2003:1012729 ANDN140:196856 TIHippocampal plasticity requires postsynaptic ephrinBs Grunwald, Ilona C.; Korte, Martin; Adelmann, Giselind; Plueck, Anne; ΑU Kullander, Klas; Adams, Ralf H.; Frotscher, Michael; Bonhoeffer, Tobias; Klein, Ruediger Department of Molecular Neurobiology, Max-Planck Institute of Neurobiology, Munich-Martinsried, 82152, Germany CS Nature Neuroscience (2004), 7(1), 33-40 CODEN: NANEFN; ISSN: 1097-6256 SO PB Nature Publishing Group DTJournal LΑ English RE.CNT THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD 38 ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 29 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. L7 STN AΝ 2004:292476 BIOSIS PREV200400291958 DN TIGenes that Influence Acute Ethanol Tolerance. Hudson, Holly R [Reprint Author]; Bhave, Sanjiv V; Tabakoff, Boris; ΑU Hoffman, Paula L Pharmacology, University of Colorado Health Sciences Center, 4200 E. Ninth Ave., Box C-236, Denver, CO, 80262, USA CS holly.hudson@uchsc.edu FASEB Journal, (2004) Vol. 18, No. 4-5, pp. Abst. 161.12. http://www.fasebj.org/. e-file. Meeting Info.: FASEB Meeting on Experimental Biology: Translating the Genome. Washington, District of Columbia, USA. April 17-21, 2004. FASEB. SO ISSN: 0892-6638 (ISSN print). DTConference; (Meeting) Conference; Abstract; (Meeting Abstract) LA English ED Entered STN: 23 Jun 2004 Last Updated on STN: 23 Jun 2004 ANSWER 30 OF 159 DISSABS COPYRIGHT (C) 2004 ProQuest Information and L7 Learning Company; All Rights Reserved on STN DISSABS Order Number: AAI3118187 2004:48564 ANEphrins and Eph receptors participate in spinal cord development and TIinjury responses in the adult

Bundesen, Liza Q. [Ph.D.]; Kromer, Lawrence F. [advisor]; Bregman, Barbara

ΑU

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CS
      Georgetown University Medical Center (0544)
      Dissertation Abstracts International, (2003) Vol. 64, No. 12B, p. 5947.
SO
      Order No.: AAI3118187. 296 pages.
DT
      Dissertation
FS
      DAI
LΑ
      English
ED
      Entered STN: 20040902
      Last Updated on STN: 20040902
L7
      ANSWER 31 OF 159
                             CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 4
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      2003:42302
                     CAPLUS
DN
      Tie-Fc and Ephrin-Fc fusion proteins for screening therapeutic capable of
TI
      modulating growth, migration and proliferation of endothelial cells and
      treating cancers
IN
      Alitalo, Kari; Kubo, Hajime
PA
      Licentia Ltd., Finland
      PCT Int. Appl., 200 pp.
SO
      CODEN: PIXXD2
DT
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LA
      English
FAN.CNT 1
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                               KIND
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                                                      APPLICATION NO.
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                                A2
      WO 2003004529
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      WO 2003004529
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      ANSWER 32 OF 159
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        2003:250468 USPATFULL
AN
        Compositions and methods for diagnosing and treating mental disorders
TI
IN
        Akil, Huda, Ann Arbor, MI, UNITED STATES
        Bunney, William E., Laguna Beach, CA, UNITED STATES
Burke, Sharon, Ann Arbor, MI, UNITED STATES
Choudary, Prabhakara V., Davis, CA, UNITED STATES
Cox, David R., Belmont, CA, UNITED STATES
Evans, Simon, Milan, MI, UNITED STATES
Jones, Edward G., Winters, CA, UNITED STATES
        Li, Jun, Palo Alto, CA, UNITED STATES
Lopez, Juan F., Ann Arbor, MI, UNITED STATES
Myers, Richard M., Stanford, CA, UNITED STATES
Thompson, Robert, Ann Arbor, MI, UNITED STATES
        Vawter, Marquis P., Laguna Niguel, CA, UNITED STATES
        Watson, Stanley J., Ann Arbor, MI, UNITED STATES
        US 2003175253
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PJ.
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        2003:225900 USPATFULL
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ΤI
        Methods for determining cell responses through EphB receptors
        Daniel, Thomas O., Nashville, TN, UNITED STATES
IN
        Stein, Elke, San Francisco, CA, UNITED STATES
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20030417 (10)
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       US 2003-420029
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           US 6555321 A 371 of International Ser. No. WO 1998-US17157, filed on
       19 Aug 1998, PENDING
       US 1997-56164P
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L7
     ANSWER 34 OF 159
AN
       2003:220740
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       Methods and compositions for diagnosing and treating rheumatoid
TI
       arthritis
       Pittman, Debra D., Windham, NH, UNITED STATES
IN
       Feldman, Jeffrey L., Arlington, MA, UNITED STATES
       Shields, Kathleen M., Harvard, MA, UNITED STATES
       Trepicchio, William L., Andover, MA, UNITED STATES
PΙ
       US 2003154032
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                                 20030814
       US 2001-23451
                            Α1
                                 20011217
ΑI
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       US 2000-255861P
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     ANSWER 35 OF 159
                        USPATFULL on STN
L7
       2003:120026
                     USPATFULL
AN
       Identification of modulatory molecules using inducible promoters Brown, Steven J., San Diego, CA, UNITED STATES
TI
IN
       Dunnington, Damien J., San Diego,
                                            CA, UNITED STATES
               Imran, San Diego, CA, UNITED STATES
       Clark,
                                 20030501
       US 2003082511
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PI
                                 20010925 (9)
ΑI
       US 2001-965201
                            Α1
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IC
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       ICS: C12Q001-68
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                        USPATFULL on STN
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AN
       2003:30255
ΤI
       B-ephrin regulation of G-protein coupled chemoattraction, compositions,
       and methods of use
       Flanagan, John G.
                           Newton, MA, UNITED STATES
IN
       Lu, Qiang, Brookline, MA, UNITED STATES
       Sun, Edna E., Brookline, MA, UNITED STATES
       US 2003022202
                                 20030130
PΙ
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       US 2002-113794
                                 20020401 (10)
AΙ
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       US 2001-280260P
                             20010330 (60)
PRAI
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IC
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        ICM: C12Q001-68
        ICS: C07H021-04; C12P021-02; C12N001-18; C12N009-16; C12N005-08
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L7
     ANSWER 37 OF 159 USPATFULL on STN
AN
        2003:115716
                       USPATFULL
ΤI
        Methods for determining cell responses through EphB receptors
        Daniel, Thomas O., Nashville, TN, United States
Stein, Elke, San Francisco, CA, United States
IN
        Vanderbilt University, Nashville, TN, United States (U.S. corporation) US 6555321 B1 20030429
PA
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PΙ
                      19990225
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                                    20000214 (9)
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        WO 1998-US17157
                                    19980819
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INCL
        INCLM: 435/007.100
        INCLS: 435/007.200; 435/007.210; 435/007.800; 435/334.000
NCL
                435/007.100
        NCLM:
                435/007.200; 435/007.210; 435/007.800; 435/334.000
        NCLS:
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        [7]
        ICM: G01N033-53
    ICS: G01N033-567; C12N005-06
435/7.1; 435/7.21; 435/7.8; 435/7.2; 435/334
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     receptor ligation.
     Yu, Guang; Luo, Hongyu; Wu, Yulian; Wu, Jiangping [Reprint Author]
Laboratory of Immunology, Research Centre, Notre Dame Hospital, CHUM,
ΑU
CS
      Sherbrooke St. East, Pavilion DeSeve, Room Y-5616, Montreal, PQ, H2L 4M1,
      Canada
      jianping.wu@umontreal.ca
     Journal of Biological Chemistry, (November 21 2003) Vol. 278, No. 47, pp.
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     Entered STN: 28 Jan 2004
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     Tabakoff, Boris [Reprint Author]; Bhave, Sanjiv V.; Hoffman, Paula L. Department of Pharmacology, University of Colorado Health Sciences Center, 4200 East Ninth Avenue, C-236, Denver, CO, 80262, USA
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     boris.tabakoff@uchsc.edu
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      Gregory; Valladeau, Jenny; Caux, Christophe; Garrone, Pierre
CS
     Laboratory for Immunological Research, Schering-Plough, 27 Chemin des
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- CS University, Sakyo-ku, Kyoto, 606-8507, Japan fuji@kuhp.kyoto-u.ac.jp Journal of Clinical Endocrinology & Metabolism, (September 2003) Vol. 88,
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- Department of Medical Biochemistry, Gothenburg University, Medicinaregatan 9 Å, 405 30, Gothenburg, Sweden CS
- klas.kullander@medkem.gu.se; ole.kiehn@neuro.ki.se Science (Washington D C), (21 March 2003) Vol. 299, No. 5614, pp. SO 1889-1892. print ISSN: 0036-8075 (ISSN print).
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- ΑU Vidovic M.; Marotte L.R.
- Dr. L.R. Marotte, Developmental Biology Group, Res. School of Biological CS Sciences, Australian National University, Canberra, ACT 0200, Australia. E-mail: marotte@rsbs.anu.edu.au
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- Matsuzawa, Yuji Department of Internal Medicine and Molecular Science, Graduate School of Modical School Osaka University, 2-2 Yamadaoka, Suita, Osaka, CS 565-0871, Japan

tamuras@imed2.med.osaka-u.ac.jp

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Rasband K; Hardy M; Chien C B (Reprint) ΑU

CS Univ Utah, Med Čtr, Dept Neurobiol & Anat, Salt Lake City, UT 84132 USA (Reprint)

CYA USA

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- DTGeneral Review; Journal

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REC Reference Count: 20

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- receptor tyrosine kinase using a scintillation proximity assay Bembenek, Michael E.; Schmidt, Stephen; Li, Ping; Morawiak, Jennifer; ΑU Prack, Andrea; Jain, Sadhana; Roy, Rebecca; Parsons, Thomas; Chee, Linda

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- CS Ophthalmology and Physiology, University of California San Francisco, San Francisco, ČĀ, USA
- SO ARVO Annual Meeting Abstract Search and Program Planner, (2003) Vol. 2003, pp. Abstract No. 5221. cd-rom.
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Conference; (Meeting Poster)

Conference; Abstract; (Meeting Abstract)

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Entered STN: 26 Nov 2003 EDLast Updated on STN: 26 Nov 2003

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 Ophthalmology and Visual Sciences, Kyoto University Graduate School of

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Medicine, Kyoto, Japan ARVO Annual Meeting Abstract Search and Program Planner, (2003) Vol. 2003, pp. Abstract No. 2881. cd-rom. SO Meeting Info.: Annual Meeting of the Association for Research in Vision and Ophthalmology. Fort Lauderdale, FL, USA. May 04-08, 2003. Association for Research in Vision and Ophthalmology.

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Conference; (Meeting Poster)

Conference; Abstract; (Meeting Abstract)

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DN PREV200400198485 TI Role of ephrins and Eph receptors in adult neurogenesis. ΑU Ricard, J. [Reprint Author]; Salinas, J. A. [Reprint Author]; Liebl, D. J. [Reprint Author] Miami Project to Cure Paralysis, Univ. of Miami, Miami, FL, USA Society for Neuroscience Abstract Viewer and Itinerary Planner, (2003) SO Vol. 2003, pp. Abstract No. 356.11. http://sfn.scholarone.com. e-file. Meeting Info.: 33rd Annual Meeting of the Society of Neuroscience. New Orleans, LA, USA. November 08-12, 2003. Society of Neuroscience. Conference; (Meeting) DT Conference; Abstract; (Meeting Abstract) LΑ English ED Entered STN: 14 Apr 2004 Last Updated on STN: 14 Apr 2004 ANSWER 61 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. L7 STN 2004:160067 $\mathbf{A}\mathbf{N}$ BIOSIS DNPREV200400160217 ΤI Corpus callosum axon guidance: the role of ephrins and Eph receptors. Mendes, S. W. [Reprint Author]; Liebl, D. J. Dept. Neurol., Univ. of Miami, Miami, FL, USA ΑU CS SO Society for Neuroscience Abstract Viewer and Itinerary Planner, (2003) Vol. 2003, pp. Abstract No. 32.17. http://sfn.scholarone.com. e-file. Meeting Info.: 33rd Annual Meeting of the Society of Neuroscience. New Orleans, LA, USA. November 08-12, 2003. Society of Neuroscience. Conference; (Meeting)
Conference; Abstract; (Meeting Abstract) DTLΑ English Entered STN: 24 Mar 2004 EDLast Updated on STN: 24 Mar 2004 ANSWER 62 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on L7 STNAN2004:159726 BIOSIS defects in synaptic functions in the hippocampus.
Rodenas-Ruano, A. I. [Reprint Author]; Liebl, D. J. [Reprint Author]; Huizinga, C. [Reprint Author]
Neurosci. Univ of Miami Vicinia. DN PREV200400159876 ΤI AU Neurosci., Univ. of Miami, Miami, FL, USA Society for Neuroscience Abstract Viewer and Itinerary Planner, CS SO Vol. 2003, pp. Abstract No. 7.4. http://sfn.scholarone.com. e-file. Meeting Info.: 33rd Annual Meeting of the Society of Neuroscience. New Orleans, LA, USA. November 08-12, 2003. Society of Neuroscience. Conference; (Meeting) Conference; Abstract; (Meeting Abstract) DT LAEnglish ED Entered STN: 24 Mar 2004 Last Updated on STN: 24 Mar 2004 L7 ANSWER 63 OF 159 DISSABS COPYRIGHT (C) 2004 ProQuest Information and Learning Company; All Rights Reserved on STN AN2003:11061 DISSABS Order Number: AAI3055126 TIAnalysis of Eph receptors and ephrin ligands in the development of hippocamposeptal projections Chen, Zhi-Yong [Ph.D.]; Zhou, Renping [adviser] ΑU Rutgers The State University of New Jersey - New Brunswick and University of Medicine and Dentistry of New Jersey (0801) CS Dissertation Abstracts International, (2002) Vol. 63, No. 5B, p. 2225. Order No.: AAI3055126. 162 pages. SO ISBN: 0-493-69309-2. DT Dissertation FS DAI LA English ANSWER 64 OF 159 IFIPAT L7 COPYRIGHT 2004 IFI on STN DUPLICATE 18 10203599 IFIPAT; IFIUDB; IFICDB ANPEPTIDES THAT MODULATE THE INTERACTION OF B CLASS EPHRINS AND PDZ TΙ DOMAINS; PEPTIDE COMPLEX FOR USE IN THE TREATMENT OF CELL PROLIFERATIVE DISORDERS IN Gish Gerald (CA); Lin Danny (CA); Pawson Anthony (CA)

Unassigned Or Assigned To Individual (68000)

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PA

PI AI US 2002147306

US 2001-862179

US 1998-109158P 19981120 (Provisional) US 2002147306 FI20021010 Utility; Patent Application - First Publication DTFS CHEMICAL APPLICATION CLMN 35 8 Figure(s). GΙ FIG. 1. Amino acid sequence of the cytoplasmic domains of the human B ephrins. Conserved residues among the three B ephrins are highlighted. Asterisks mark conserved tyrosines that are potential sites of phosphorylation. The potential PDZ domain binding site is underlined. FIGS. 2A-D. Identification of PDZ domain-containing candidates for ephrin B binding. FIG. 2A, The preferred binding sequence of FAP-1 PDZ5 is shown below a schematic representation of the entire FAP-1 protein tyrosine below a schematic representation of the entire FAP-1 protein tyrosine phosphatase. FAP-1 PDZ5 domain specificity was deduced from an oriented peptide library technique (1). Residues within the optimal binding sequence that match the C-terminal sequence of B ephrins are indicated in bold. The organization of the PDZ domains of FAP-1 shown in this figure follows the numbering described by Sato et al. (33). FIG. 2B, Diagrammatic representations of the PDZ domaincontaining proteins identified through an expression screen with a biotinylated peptide probe of eprhin B3 C-terminal sequence. The brackets mark the portions of the protein encoded by the cDNAs isolated from the screen. PDZ domains are protein encoded by the cDNAs isolated from the screen. PDZ domains are represented by grey boxes. FIG. 2C, Amino acid sequence alignment of FAP-1 PDZ5 and of the PDZ domains isolated in the expression screen. The numbering of the PDZ domains is as shown in FIG. 2B. Conserved residues are highlighted. The alignment was performed with the ClustalW program (55). FIG. 2D, Amino acid sequence alignment of PHIP and PAR-3. Conserved residues are highlighted and the PDZ domains are underlined. The alignment was performed with the Genestream Align program.

FIGS. 3A-C. FAP-1 PDZ5 and syntenin bind specifically to ephrin B1 in GST-mixes. Cos-1 cells were transiently transfected with either wild-type ephrin B1 (W. T.) or the ephrin B1 Val deletion (Val Delta) or were untransfected. Cell lysates were incubated with the GST fusion proteins as indicated and analyzed by immunoblotting with anti-ephrin B1 antibody. untransfected. Cell lysates were incubated with the GST fusion proteins as indicated and analyzed by immunoblotting with anti-ephrin B1 antibody. Immunoprecipitated ephrin B1 or ephrin B1 Val Delta were included as a positive control. FIG. 3A and FIG. 3B, GST-mixes with fusion proteins of FAP-1. C and D, GST-mixes with fusion proteins of syntenin.

FIGS. 4A and 4B. FAP-1 PDZ5 and syntenin binding to ephrin B1 can be blocked by addition of peptides corresponding to the Cterminal sequence of B ephrins. Peptides of the indicated sequence were included at a concentration of 100 mu M in incubations of GST fusion proteins with lysates of Cos-1 cells transfected with ephrin B1. Associated proteins were separated on a 10% polyacrylamide/SDS gel and analyzed by immunoblotting with antibodies against ephrin B1. FIG. 4A, Competition of FAP1 PDZ5 binding to ephrin B1 using the indicated peptides. A peptide of FAP1 PDZ5 binding to ephrin B1 using the indicated peptides. A peptide of sequence DHQpYpYND was added at a concentration of 100 mu M as a negative control. Immunoprecipitation of ephrin B1 was included as a positive control. FIG. 4B, Peptide competition of the binding of full-length syntenin to ephrin B1.

FIGS. 5A and 5B. Fluorescence polarization analysis of GST-FAP-1 PDZ3, GST-FAP-1 PDZ5 and GST-syntenin binding to Fluoresceinlabelled peptides corresponding to the C-terminus of ephrin B1. FIG. 5 A, Solutions containing the indicated final concentration of GST-FAP-1 PDZ3 (FIG. 5B) containing the indicated final concentration of GST-FAP-1 PDZ3 (FIG. 5B, A Binding of a GST fusion of full-length syntenin to the NIYYKV (FIG. 6. Co-immunoprecipitation of syntenin-FLAG with ephrin B1. Cos-1 cells were co-transfected with either ephrin B1 and syntenin-FLAG or with the ephrin B1 Val deletion and synteninFLAG as indicated. Cell lysates were immunoprecipitated with antibodies against ephrin B1 or IL-3 receptor a or were treated with protein A sepharose only. Immunocomplexes were subjected to SDS-PAGE (10%) and blotted with anti-FLAG antibodies. FIG. 7. Fluorescence polarization analysis of GST-PHIP PDZ3 binding to Fluorescein-labelled peptides corresponding to the Cterminus of ephrin B1. Solutions containing the indicated final concentration of GST-PHIP PDZ3 fusion protein in mixtures containing 25 nM fluorescein-labelled peptide probe, 20 mM phosphate pH 7.0, 100 mM NaCl, and 2 mM DTT were monitored for fluorescence polarization at 22 degrees C. The GST-PHIP PDZ3 fusion protein was measured for binding to the phosphorylated PDZ3 fusion protein was measured for binding to the phosphorylated peptides, NIPYYKV (), NiYpYKV () and NIPYPYKV (
FIG. 8 PHIP PDZ3 binds specifically to V-Src phosphorylated ephrin B1 in
GST-mixes. COS-1 cells were transiently cotransfected with V-Src and
either wild-type ephrin B1 or the ephrin B1 Val deletion (VA) or were
transfected with either wild-type ephrin B1 or ephrin B1 Val deletion
alone. Cell lysates were incubated with the GST fusion proteins as
indicated and analyzed by immunoblotting with antiphosphotyrosine

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       London, WC1 6BT, UK
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       CODEN: ESRWEL; ISSN: 0947-6075
PB
       Springer-Verlag
       Journal; General Review
DT
LΑ
       English
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                   THERE ARE 81 CITED REFERENCES AVAILABLE FOR THIS RECORD
                   ALL CITATIONS AVAILABLE IN THE RE FORMAT
L7
       ANSWER 100 OF 159
                                            COPYRIGHT (c) 2004 The Thomson Corporation.
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                                                                                                            on
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       STN
AN
       2000:93783
                      BIOSIS
DN
       PREV200000093783
       Expression of EphA4 in developing inner ears of the mouse and guinea pig. van Heumen, Walter R. A.; Claxton, Christina; Pickles, James O. [Reprint
TI
ΑU
       author]
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Vision, Touch and Hearing Research Centre, Department of Physiology and Pharmacology, University of Queensland, Brisbane, QLD, 4072, Australia Hearing Research, (Jan., 2000) Vol. 139, No. 1-2, pp. 42-50. print.

CS

SO

CODEN: HERED3. ISSN: 0378-5955.

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LΑ
      English
ED
      Entered STN: 10 Mar 2000
      Last Updated on STN: 3 Jan 2002
      ANSWER 101 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation.
L7
      STN
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      2001:87122
                    BIOSIS
DN
      PREV200100087122
TI
      Aberrant retinocollicular mapping in EphB2/EphB3 double knockout mice.
ΑU
      Hindges, R. [Reprint author]; McLaughlin, T.; Henkemeyer, M.; O'Leary, D.
CS
      The Salk Institute, La Jolla, CA, USA
      Society for Neuroscience Abstracts, (2000) Vol. 26, No. 1-2, pp. Abstract
SO
      No.-218.1. print.
      Meeting Info.: 30th Annual Meeting of the Society of Neuroscience. New Orleans, LA, USA. November 04-09, 2000. Society for Neuroscience.
      ISSN: 0190-5295.
DT
      Conference; (Meeting)
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LA
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      Entered STN: 14 Feb 2001
ED
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      ANSWER 102 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation.
L7
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      2001:87002 BIOSIS
AN
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TT
      Distinct functions of EphA and EphB receptors in hippocamposeptal
      topographic map formation.
AU
      Chen, Z. [Reprint author]; Yue, Y.; Su, J.; Sun, C.; Henkemeyer, M.; Zhou,
CS
      Rutgers University, Piscataway, NJ, USA
SO
      Society for Neuroscience Abstracts, (2000) Vol. 26, No. 1-2, pp. Abstract
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      ANSWER 103 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation.
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      2001:96762 BIOSIS
AN
      PREV200100096762
DN
                         ***ephrin*** - ***B3***
ΤI
      Ephrin-B2 and
                                                           expression in the inner ear.
      Blanchi, L. M. [Reprint author]; Dinsio, K.; Gale, N. W.; Henkemeyer, M.;
AU
      Fritzsch, B.
CS
      Oberlin College, Oberlin, OH, USA
SO
      Society for Neuroscience Abstracts, (2000) Vol. 26, No. 1-2, pp. Abstract
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      Meeting Info.: 30th Annual Meeting of the Society of Neuroscience. New Orleans, LA, USA. November 04-09, 2000. Society for Neuroscience.
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      Entered STN: 21 Feb 2001
      Last Updated on STN: 15 Feb 2002
      ANSWER 104 OF 159 DISSABS COPYRIGHT (C) 2004 ProQuest Information and
L7
      Learning Company; All Rights Reserved on STN
      2000:8846 DISSABS Order Number: AAIMQ42194
IDENTIFICATION OF DIFFERENTIALLY EXPRESSED GENES DURING DIFFERENTIATION OF A NOVEL HUMAN VASCULAR SMOOTH MUSCLE CELL LINE
PRAVDA, ZUZANA HEDRIKA [M.SC.]; PICKERING, J. G. [adviser]
THE UNIVERSITY OF WESTERN ONTARIO (CANADA) (0784)
AN
TI
ΑU
CS
      Masters Abstracts International, (1999) Vol. 38, No. 1, p. 163. Order No.:
SO
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AAIMQ42194. 150 pages. ISBN: 0-612-42194-5.

Dissertation

MAI

English

DT

FS

LΑ

ANSWER 105 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. L7 STN DUPLICATE 32 1999:134203 BIOSIS ΔN DN PREV199900134203 The carboxyl terminus of B class ephrins constitutes a PDZ domain binding ΤI motif. Lin, Dan; Gish, Gerald D.; Songyang, Zhou; Pawson, Tony [Reprint author] Programmne Mol. Biol. Cancer, Samuel Lunenfeld Res. Inst., Mt. Sinai Hosp., 600 University Ave., Toronto, ON M5G 1X5, Canada Journal of Biological Chemistry, (Feb. 5, 1999) Vol. 274, No. 6, pp. AU CS SO 3726-3733. print. CODEN: JBCHĀ3. ISSN: 0021-9258. DT Article LA English Entered STN: 31 Mar 1999 ED Last Updated on STN: 31 Mar 1999 ANSWER 106 OF 159 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation. L7 on STN 1999:975353 SCISEARCH ANThe Genuine Article (R) Number: 250YD GA ***EphrinB3*** and EphB3 are coordinately upregulated during human ΤI smooth muscle cell maturation ΑU Pravda Z (Reprint); Li S; Rajakumar N; Ruschlow W; Verdi J; Brown A; Pickering J G JOHN P RÕBARTS RES INST, LONDON, ON N6A 5K8, CANADA; UNIV WESTERN ONTARIO, CS LONDON, ON, CANADA CYA CANADA CIRCULATION, (2 NOV 1999) Vol. 100, No. 18, Supp. [S], pp. 3636-3636. Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA SO 19106-3621. ISSN: 0009-7322. DTConference; Journal FS LIFE; CLIN LA English REC Reference Count: 0 L7 ANSWER 107 OF 159 CANCERLIT on STN DUPLICATE 33 CANCERLIT AN1999316799 99316799 PubMed ID: 10389937 DN High-level expression of EPHB6, EFNB2, and EFNB3 is associated with low TI tumor stage and high TrkA expression in human neuroblastomas. Tang X X; Evans A E; Zhao H; Cnaan A; London W; Cohn S L; Brodeur G M; ΑU Ikegaki N CS Division of Oncology, The Children's Hospital of Philadelphia, Abramson Research Center, Pennsylvania 19104-4318, USA. CA70958 (NCI) F32 CA75748 (NCI) NC NS34514 (NINDS) SO CLINICAL CANCER RESEARCH, (1999 Jun) 5 (6) 1491-6. Journal code: 9502500. ISSN: 1078-0432. CY United States DTJournal; Article; (JOURNAL ARTICLE) LА MEDLINE; Priority Journals MEDLINE 1999316799 FS OS EM199909 ED Entered STN: 19991112 Last Updated on STN: 19991112 L7 ANSWER 108 OF 159 CAPLUS COPYRIGHT 2004 ACS on STN AN1999:223544 CAPLUS 131:30116 DN TI EphrinB ligands recruit GRIP family PDZ adaptor proteins into raft membrane microdomains Bruckner, Katja; Labrador, Juan Pablo; Scheiffele, Peter; Herb, Anne; Seeburg, Peter H.; Klein, Rudiger AU Developmental Biology Programme, European Molecular Biology Laboratory, Heidelberg, D-69117, Germany Neuron (1999), 22(3), 511-524 CODEN: NERNET; ISSN: 0896-6273 CS SO Cell Press PB DT Journal LAEnglish RE.CNT THERE ARE 54 CITED REFERENCES AVAILABLE FOR THIS RECORD

L7 ANSWER 109 OF 159 CANCERLIT on STN DUPLICATE 34 ΑN 1999154801 CANCERLIT DN 99154801 PubMed ID: 10037197 Coexpression of transcripts encoding EPHB receptor protein tyrosine kinases and their ephrin-B ligands in human small cell lung carcinoma. Tang X X; Brodeur G M; Campling B G; Ikegaki N Division of Oncology, The Children's Hospital of Philadelphia, Pennsylvania 19104-4318, USA. TI AII CS NC CA70958 (NCI) F32 CA75748 (NCI) NS34514 (NINDS) SO CLINICAL CANCER RESEARCH, (1999 Feb) 5 (2) 455-60. Journal code: 9502500. ISSN: 1078-0432. CY United States DT Journal; Article; (JOURNAL ARTICLE) LΑ English MEDLINE; Priority Journals MEDLINE 1999154801 FS OS EM 199904 ED Entered STN: 19990622 Last Updated on STN: 19990622 ANSWER 110 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. L7 STN DUPLICATE 35 AN 2000:51186 BIOSIS PREV20000051186 DN Comparative analysis of embryonic gene expression defines potential TIinteraction sites for Xenopus EphB4 receptors with ephrin-B ligands. Helbling, Paul M.; Saulnier, Didier M.E.; Robinson, Vicky; Christiansen, ΑU Jeff H.; Wilkinson, David G.; Brandli, Andre W. [Reprint author] Institute of Cell Biology, Swiss Federal Institute of Technology, ETH-Honggerberg, CH-8093, Zurich, Switzerland Developmental Dynamics, (Dec., 1999) Vol. 216, No. 4-5, pp. 361-373. CS SO print. CODEN: DEDYEI. ISSN: 1058-8388. DTArticle LAEnglish ED Entered STN: 3 Feb 2000 Last Updated on STN: 3 Jan 2002 ANSWER 111 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. L7 STNNA 1999:143675 BIOSIS DNPREV199900143675 Roles of ephrinB ligands and EphB receptors in cardiovascular development: Demarcation of arterial/ venous domains, vascular morphogenesis, and TI sprouting angiogenesis.
Adams, Ralf H.; Wilkinson, George A.; Weiss, Cornelia; Diella, Francesca; AU Gale, Nicholas W.; Deutsch, Urban; Risau, Werner; Klein, Ruediger [Reprint author] European Mol. Biol. Lab., D-69117 Heidelberg, Germany CS SO Genes and Development, (Feb. 1, 1999) Vol. 13, No. 3, pp. 295-306. print. CODEN: GEDEEP. ISSN: 0890-9369. DT Article LΑ English Entered STN: 31 Mar 1999 Last Updated on STN: 31 Mar 1999 ED L7 ANSWER 112 OF 159 MEDLINE on STN AN1999210443 MEDLINE DNPubMed ID: 10192794 TI Induction of Eph B3 after spinal cord injury. Miranda J D; White L A; Marcillo A E; Willson C A; Jagid J; Whittemore S R Department of Neurological Surgery, University of Miami School of ΑU CS Medicine, 1600 Northwest 10th Avenue, R-48, Miami, Florida 33136, USA. NS10304 (NINDS) NC NS26887 (NINDS) Experimental neurology, (1999 Mar) 156 (1) 218-22. Journal code: 0370712. ISSN: 0014-4886. SO United States CY DT Journal; Article; (JOURNAL ARTICLE) LA English FS Priority Journals 199904

EΜ

Last Updated on STN: 20000303 Entered Medline: 19990421 L7 ANSWER 113 OF 159 MEDLINE on STN 1999172312 ANMEDLINE PubMed ID: 10072375 DNTIEph receptors and ephrins in neural development. ΑU O'Leary D D; Wilkinson D G Molecular Neurobiology Laboratory The Salk Institute 10010 North Torrey CS Pines Road La Jolla Čalifornia 92037 USA.. dennis oleary@qm.salk.edu NC (NEI) NS31558 (NINDS) Current opinion in neurobiology, (1999 Feb) 9 (1) 65-73. Ref: 74 SO Journal code: 9111376. ISSN: 0959-4388. CYENGLAND: United Kingdom Journal; Article; (JOURNAL ARTICLE) General Review; (REVIEW) (REVIEW LITERATURE) DT LΑ English FS Priority Journals 199905 EMEntered STN: 19990607 EDLast Updated on STN: 19990607 Entered Medline: 19990524 L7 ANSWER 114 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. STN AN2000:25109 BIOSIS PREV200000025109 DN***EphrinB3*** and EphB3 are coordinately upregulated during human TIsmooth muscle cell maturation. Pravda, Zuzana [Reprint author]; Li, Shaohua [Reprint author]; Rajakumar, ΑU N.; Ruschlow, Walter; Verdi, Joseph; Brown, Arthur; Pickering, J. Geoffrey John P Robarts Res Inst, London, ON, Canada Circulation, (Nov. 2, 1999) Vol. 100, No. 18 SUPPL., pp. I.689. print. Meeting Info.: 72nd Scientific Sessions of the American Heart Association. CS SO Atlanta, Georgia, USA. November 7-10, 1999. CODEN: CIRCAZ. ISSN: 0009-7322. Conference; (Meeting) DT Conference; Abstract; (Meeting Abstract) LΑ English EDEntered STN: 29 Dec 1999 Last Updated on STN: 31 Dec 2001 ANSWER 115 OF 159 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. L7 STN AN1999:64189 BIOSIS DNPREV199900064189 Induction of EPH B3 RPTK after spinal cord injury. TIMiranda, J. D.; White, L. A.; Willson, C. A.; Marcillo, A.; Jagid, J.; AU Whittemore, S. R. CS Miami Project Dep. Neurolgoical Surgery, Univ. Miami Sch. Med., Miami, FL 33136, USĀ Society for Neuroscience Abstracts, (1998) Vol. 24, No. 1-2, pp. 741. SO print. Meeting Info.: 28th Annual Meeting of the Society for Neuroscience, Part 1. Los Angeles, California, USA. November 7-12, 1998. Society for Neuroscience. ISSN: 0190-5295. DT Conference; (Meeting) Conference; Abstract; (Meeting Abstract) Conference; (Meeting Poster)

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Bergemann, Andrew D.; Zhang, Lee; Chiang, Ming-Ko; Brambilla, Riccardo;

Dep. Cell Biol., Harv. Med. Sch., 240 Longwood Ave., Boston, MA 02115, USA

on

DUPLICATE 36

a ligand for the receptor EphB3, expressed

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CS

English

STN

Entered STN: 16 Feb 1999

1998:132637 BIOSIS

PREV199800132637

Ephrin

Last Updated on STN: 16 Feb 1999

B3

Klein, Ruediger; Flanagan, John G. [Reprint author]

at the midline of the developing neural tube.

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CODEN: ONCNES. ISSN: 0950-9232.
DT
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     Entered STN: 20 Mar 1998
     Last Updated on STN: 20 Mar 1998
L7
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                             MEDLINE on STN
     1998191577
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AN
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DN
TI
     The ephrins and Eph receptors in neural development.
     Flanagan J G; Vanderhaeghen P
Department of Cell Biology, Harvard Medical School, Boston, Massachusetts
ΑU
CS
     02115, USA.. Flanagan@warren.med.harvard.edu
SO
     Annual review of neuroscience, (1998) 21 309-45. Ref: 185
     Journal code: 7804039. ISSN: 0147-006X.
CY
     United States
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General Review; (REVIEW)
DT
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AN
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DN
     Loss of EphB2 and B3 receptor tyrosine kinases results in pathfinding
TI
     errors of retinal ganglion cell axons to the optic disc.
     Birgbauer, E. [Reprint author]; Henkemeyer, M.; Sretavan, D. [Reprint
ΑU
     U.C. San Francisco, San Francisco, CA, USA
CS
SO
     Molecular Biology of the Cell, (Nov., 1998) Vol. 9, No. SUPPL., pp. 228A.
     print.
     Meeting Info.: 38th Annual Meeting of the American Society for Cell
     Biology. San Francisco, California, USA. December 12-16, 1998. American Society for Cell Biology.
     CODEN: MBCEEV. ISSN: 1059-1524.
     Conference; (Meeting)
Conference; Abstract; (Meeting Abstract)
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     Entered STN: 20 Jan 1999
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     ANSWER 119 OF 159
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     PubMed ID: 9126477
DN
     cDNA cloning, chromosomal localization, and expression pattern of EPLG8, a new member of the EPLG gene family encoding ligands of EPH-related
TI
     protein-tyrosine kinase receptors.
ΑU
     Tang X X; Pleasure D E; Ikegaki N
CS
     Division of Neurology Research, Children's Hospital of Philadelphia,
     Pennsylvania 19104-4318, USA.
NC
     NS08075 (NINDS)
     NS25044 (NINDS)
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                (1997 Apr 1) 41 (1) 17-24.
     Genomics,
     Journal code: 8800135. ISSN: 0888-7543.
CY
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     Journal; Article; (JOURNAL ARTICLE)
DT
LА
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L7
      ANSWER 120 OF 159 DGENE
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AN
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      Identifying modulators of binding between a Tie receptor tyrosine kinase
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IN
       Alitalo K; Kubo H
PA
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PΙ
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DESC
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L7
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TI
       Identifying modulators of binding between a Tie receptor tyrosine kinase
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       Isolated complex for treating proliferative or differentiative disorders
TI
       comprises B class ephrin and PDZ domain containing protein -
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       Lin D; Pawson A
                     MOUNT SINAI HOSPITAL.
PA
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PΙ
      WO 2000031124 A2 20000602
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      WO 1999-CA1101
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      ANSWER 123 OF 159
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OS
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DESC
      EphrinB-3 C-terminal peptide biotinylated probe, comprising PDZ domain.
L7
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      ANSWER 124 OF 159
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      Use of a compound or composition for diagnosing, treating or preventing synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
TI
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IN
      Nakamura Y; Kataqiri T
                     ONCOTHERAPY SCI INC.
PA
       (ONCO-N)
       (UYTY)
                     UNIV TOKYO.
PΙ
      WO 2004020668 A2 20040311
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DESC
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                                     COPYRIGHT 2004 The Thomson Corp on STN
L7
      ANSWER 125 OF 159
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ΤI
       Use of a compound or composition for diagnosing, treating or preventing
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IN
       Nakamura Y; Kataqiri T
                    ONCOTHERAPY SCI INC.
PA
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       (UYTY)
                    UNIV TOKYO
PΙ
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DESC
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      ANSWER 126 OF 159
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      Use of a compound or composition for diagnosing, treating or preventing
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      myeloid leukemia.
TN
      Nakamura Y; Katagiri T
PA
                    ONCOTHERAPY SCI INC.
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       (UYTY)
                    UNIV TOKYO
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L7
      ANSWER 127 OF 159
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IN
      Nakamura Y; Katagiri T
PA
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L7
      ANSWER 128 OF 159
                            DGENE
                                   COPYRIGHT 2004 The Thomson Corp on STN
AN
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TI
      Use of a compound or composition for diagnosing, treating or preventing synovial sarcoma or a disease associated with Frizzled homologue 10, e.e.
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IN
      Nakamura Y; Katagiri T
PA
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OS
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      ANSWER 129 OF 159
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L7
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TI
       Use of a compound or composition for diagnosing, treating or preventing
       synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
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       Nakamura Y; Katagiri T
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PA
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L7
      ANSWER 130 OF 159
                            DGENE
                                   COPYRIGHT 2004 The Thomson Corp on STN
      ADL91707
AN
                 DNA
                             DGENE
ΤI
      Use of a compound or composition for diagnosing,
                                                             treating or preventing
       synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
       colorectal cancer, gastric cancer, chronic myeloid leukemia or acute
      myeloid leukemia.
IN
      Nakamura Y; Katagiri T
PA
       (ONCO-N)
                    ONCOTHERAPY SCI INC.
       (UYTY)
                    UNIV TOKYO
PΙ
      WO 2004020668 A2 20040311
                                                   143p
ΑI
      WO 2003-JP10591
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      US
         2002-407506P
PRAI
                              20020830
      US 2003-486195P
                              20030711
DT
      Patent
LΑ
      English
OS
      2004-239208 [22]
DESC
         ***Ephrin***
                            ***B3***
                                         (SYX 8) control S-oligonucleotide, SEQ ID
      NO:108.
                                   COPYRIGHT 2004 The Thomson Corp on STN
L7
      ANSWER 131 OF 159
                            DGENE
AN
      ADL91705
                 DNA
                             DGENE
TT
      Use of a compound or composition for diagnosing, treating or preventing synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
      colorectal cancer, gastric cancer, chronic myeloid leukemia or acute
      myeloid leukemia.
IN
      Nakamura Y; Katagiri T
PA
                    ONCOTHERAPY SCI INC.
       (ONCO-N)
       (UYTY)
                    UNIV TOKYO
PI
      WO 2004020668 A2 20040311
                                                   143p
AΙ
      WO 2003-JP10591
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DT
      Patent
LΑ
      English
OS
      2004-239208 [22]
DESC
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      NO:106.
L7
      ANSWER 132 OF 159
                           DGENE
                                   COPYRIGHT 2004 The Thomson Corp on STN
AN
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                 DNA
TI
      Use of a compound or composition for diagnosing, treating or preventing synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
      colorectal cancer, gastric cancer, chronic myeloid leukemia or acute
      myeloid leukemia.
IN
      Nakamura Y; Katagiri T
PA
       (ONCO-N)
                    ONCOTHERAPY SCI INC.
       (UYTY)
                    UNIV TOKYO
PI
      WO 2004020668 A2 20040311
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AΙ
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PRAI
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DT
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LΑ
      English
OS
      2004-239208 [22]
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DESC
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      NO:102.
                                   COPYRIGHT 2004 The Thomson Corp on STN
L7
      ANSWER 133 OF 159
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Use of a compound or composition for diagnosing, treating or preventing synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
TI
       colorectal cancer, gastric cancer, chronic myeloid leukemia or acute
       myeloid leukemia.
IN
       Nakamura Y; Katagiri T
PA
                    ONCOTHERAPY SCI INC.
       (ONCO-N)
                    UNIV TOKYO.
       (YTYU)
PΙ
       WO 2004020668 A2 20040311
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ΑI
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       US 2003-486195P
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OS
       2004-239208 [22]
DESC
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L7
       ANSWER 134 OF 159
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                                    COPYRIGHT 2004 The Thomson Corp on STN
AN
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                             DGENE
TI
       Use of a compound or composition for diagnosing, treating or preventing
       synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
       colorectal cancer, gastric cancer, chronic myeloid leukemia or acute
       myeloid leukemia.
IN
       Nakamura Y; Katagiri T
PA
                    ONCOTHERAPY SCI INC.
       (ONCO-N)
       (UYTY)
                    UNIV TOKYO.
       WO 2004020668 A2 20040311
WO 2003-JP10591 2003
PI
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DT
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       English
LΑ
OS
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DESC
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                                         (SYX 8) control S-oligonucleotide, SEQ ID
       NO:105.
L7
       ANSWER 135 OF 159
                            DGENE
                                    COPYRIGHT 2004 The Thomson Corp on STN
AN
       ADL91700
                 DNA
                             DGENE
TI
       Use of a compound or composition for diagnosing, treating or preventing synovial sarcoma or a disease associated with Frizzled homologue 10, e.g.
       colorectal cancer, gastric cancer, chronic myeloid leukemia or acute
       myeloid leukemia.
IN
       Nakamura Y; Katagiri T
PA
       (ONCO-N)
                    ONCOTHERAPY SCI INC.
       (UYTY)
                    UNIV TOKYO.
      WO 2004020668 A2 20040311
WO 2003-JP10591 2003
US 2002-407506P 2002
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PRAI
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DT
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LΑ
       English
OS
       2004-239208 [22]
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DESC
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L7
                                    COPYRIGHT 2004 The Thomson Corp on STN
       ANSWER 136 OF 159
                            DGENE
AN
       ABX12547
                 CDNA
                              DGENE
ΤI
       Identifying modulators of binding between a Tie receptor tyrosine kinase
       and an Ephrin_ligand, useful for promoting neovascularization, comprises
       contacting a Tie receptor with an Ephrin in the presence of a putative
       modulator
IN
       Alitalo K; Kubo H
PA
       (LICN)
                    LICENTIA LTD.
ΡI
       WO 2003004529 A2 20030116
                                                    199p
                              20020702
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       US 2001-302960P
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       2003-210341 [20]
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CR
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DESC
       cDNA encoding mouse
                                ***ephrin*** - ***B3***
L7
                            DGENE
       ANSWER 137 OF 159
                                    COPYRIGHT 2004 The Thomson Corp on STN
ΑN
       ABX12546
                 CDNA
                              DGENE
       Identifying modulators of binding between a Tie receptor tyrosine kinase
TI
       and an Ephrin ligand, useful for promoting neovascularization, comprises
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modulator -
IN
         Alitalo K; Kubo H
PA
         (LICN)
                          LICENTIA LTD.
        WO 2003004529 A2 20030116
WO 2002-IB2524 2002
US 2001-302960P 2001
PΙ
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PRAI
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         2003-210341 [20]
         P-PSDB: ABU07845
CR
                                        ***ephrin*** - ***B3***
DESC
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                                                                                ligand.
L7
         ANSWER 138 OF 159
                                  DGENE
                                              COPYRIGHT 2004 The Thomson Corp on STN
AN
         AAA52948
                      DNA
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         Method of facilitating regeneration, growth and/or development of a central nervous system in an animal or bird, for treating disease or
TI
        trauma comprises modifying levels of Eph receptor -
Bartlett P F; Hartley L; Pilizzotto M; Kilpatrick T; Kontgen F; Coonan J;
Greferath U; Boyd A W; Dottori M; Galea M; Paxinos G; Murphy M
(HALL-N) HALL INST MEDICAL RES WALTER & ELIZA.
IN
PA
         (COUN-N)
                          COUNCIL QUEENSLAND INST MEDICAL RES.
                          UNIV MELBOURNE.
PΙ
         WO 2000024413 A1 20000504
                                                                   48p
         WO 1999-AU931
                                       19991027
ΑI
        AU 1998-6748
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PRAI
DT
         Patent
         English
LΑ
         2000-350585 [30]
OS
DESC
                   ***Ephrin***
                                          ***B3***
        Mouse
                                                            cDNA PCR primer #2.
L7
        ANSWER 139 OF 159 DGENE
                                             COPYRIGHT 2004 The Thomson Corp on STN
AN
                      DNA
                                     DGENE
        Method of facilitating regeneration, growth and/or development of a central nervous system in an animal or bird, for treating disease or
TI
        trauma comprises modifying levels of Eph receptor -
Bartlett P F; Hartley L; Pilizzotto M; Kilpatrick T; Kontgen F; Coonan J;
Greferath U; Boyd A W; Dottori M; Galea M; Paxinos G; Murphy M
(HALL-N) HALL INST MEDICAL RES WALTER & ELIZA.
IN
PA
                          COUNCIL QUEENSLAND INST MEDICAL RES.
         (COUN-N)
                          UNIV MELBOURNE.
         (UYME)
PΙ
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                                                                   48p
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                                      19981027
DT
        Patent
LΑ
        English
         2000-350585 [30]
OS
DESC
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        Mouse
                    ***Ephrin***
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L7
       ANSWER 140 OF 159
                                  FEDRIP COPYRIGHT 2004 NTIS on STN
AN
       2004:191375
                         FEDRIP
NR
       CRISP 2P01HD23315-16
                                         0006
ΤI
       EPH FAMILY RECEPTORS & LIGANDS
                                                     IN HIPPOCAMPAL SYSTEM
SF
       Principal Investigator: ZHOU, RENPING; UMDNJ-ROBERT W JOHNSON MED SCH, 675
       HOES LANE
CSP
CSS
       UNIV OF MED/DENT NJ-R W JOHNSON MED SCH, PISCATAWAY, NEW JERSEY
       Supported By: NATIONAL INSTITUTE OF CHILD HEALTH AND HUMAN DEVELOPMENT
              (/01/87)
DB
       2007
FYR
       2003
DE
       2003 (/31/08)
       Competing Continuation (Type 2)
FU
FS
       National Institutes of Health
L7
       ANSWER 141 OF 159
                                 FEDRIP COPYRIGHT 2004 NTIS on STN
AN
       2004:164868
                         FEDRIP
NR
       CRISP 5R01CA85519-03
       Functions of EPH Receptors and Ephrins in Neuroblastoma
Principal Investigator: IKEGAKI, NAOHIKO; IKEGAKI@EMAIL.CHOP.EDU,
CHILDREN'S HOSPITAL OF PHILADELP, 3516 CIVIC CENTER BLVD
CHILDREN'S HOSPITAL OF PHILADELPHIA, PHILADELPHIA, PENNSYLVANIA
Supported By: NATIONAL CANCER INSTITUTE
2007 (/01/01)
TI
SF
CSP
CSS
DB
FYR
       2003
DE
       2006 (/30/06)
       Noncompeting Continuation (Type 5)
FU
FS
       National Institutes of Health
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ANSWER 142 OF 159
L7
                                   GENBANK.RTM.
                                                      COPYRIGHT 2004 on STN
LOCUS (LOC): BV209208
GenBank ACC. NO. (GBN): BV209208
GenBank VERSION (VER): BV209208
                                                  GenBank (R)
                                 BV209208.1 GI:51853752
CAS REGISTRY NO. (RN): SEQUENCE LENGTH (SQL):
                                 753950-77-5
                                 706
MOLECULE TYPE (CI):
                                 DNA; linear
DIVISION CODE (CI):
                                 Sequence Tag Site
                                2 Sep 2004
EFNB3_3177 Rhesus macaque genomic DNA Macaca mulatta
DATE (DATE):
DEFINITION (DEF):
                                STS genomic clone MMA3177, sequence tagged site.
KEYWORDS (ST):
                                STS
SOURCE:
                                Macaca mulatta (rhesus monkey)
 ORGANISM (ORGN):
                                Macaca mulatta
                                 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                                Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                                Cercopithecidae; Cercopithecinae; Macaca
COMMENT:
      Contact: Spindel ER
      Division of Neuroscience
      Oregon National Primate Research Center
       505 NW 185th Avenue, Beaverton, OR 97006, USA
       Tel: 403-690-5388
      Fax: 503-690-5384
      Email: spindele@ohsu.edu
Primer A: gactgtaagaggttagaggtg
Primer B: aattcaagtccagtcattct
      STS size: 706
      PCR Profile:
              Hot Start: 95 degrees C for 2.00 min
Denaturation: 95 degrees C for 0.50 min
Annealing: 51 doggees C for 0.50 min
                                     51 degrees C for 0.50 min
               Annealing:
                                     72 degrees C for 1.00 min
               Polymerization:
               PCR Cycles:
                                      35
              Extension 72 degrees C for 7.0 min Thermal Cycler: MJ Instruments PTC100
      Protocol:
                      Template:
                                            200 ng
                     Primer:
                                            each luM
                     dNTP's:
                                            each 200 uM
                     Taq Polymerase: 0.05 units/ul (Fast Start High
      Fidelity, Roche)
                     Total Vol:
                                           50 ul
      Buffer:
                    MqCl2:
                                           1.8 mM
      Fast Start polymerase reaction buffer (Roche)
Bases 1-697 are 95% homologous (Blast) to bases 2529-3219 of
NM 001406.2. Primers were chosen to amplify genomic DNA in the 3'
region of EFNB3. As human sequence was used to design the primers,
the primer sequences are not included in the resulting sequence
      provided below. To obtain additional information regarding
      primers or clones contact: Dr. Robert Norgren; Dept of Genetics,
      Cell Biology & Anatomy; University of Nebraska Medical Center; 986395 Nebraska Medical Center; Omaha, NE 68198. Email:
      rnorgren@unmc.edu
      A database containing sequences associated with this project can be found at: http://rhesusgenechip.unomaha.edu/index.html.
                                1 (bases 1 to 706)
Spindel, E.R.; Pauley, M.; Jia, Y.; Boyle, N.; Jiang, S.;
Gravett, C.; Lupo, S.L.; Ali, H.; Ojeda, S.R.; Norgren, R.B.
Targeted amplification of the 3' end of rhesus macaque
REFERENCE:
    AUTHOR (AU):
    TITLE (TI):
                                orthologs of human genes
    JOURNAL (SO):
                                Unpublished (2004)
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                          Location
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DNA"

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      181 gaatčeatgt gtteteeega gtaacečaga tggetgtett gtteatteča teeteeattt
     241 ccgactcett tcagactcaa catagttccc ttcttagtga ccaaaatggt ggcctactgg
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421 ttttattaga tcaaagctgc tgttgggcac caggtcggcc acctcaatca ccagccaaga
481 tggttgcttt gtccaccaga ggtcaatcac ctctctggt ctgtagttcc cagctccttc
541 ctgatttttc tacaccaga ggtcaatcac caggaagttg atattgccat gggggg
      601 tatgccgtca cctcaatagt tttactgtaa aagggaaatt tgaacaacaa aaaccaaaaa
      661 aataaaaata aaaataaaaa acttcaaaag ttaacaagaa ggctgg
L7
         ANSWER 143 OF 159
                                           GENBANK.RTM.
                                                                          COPYRIGHT 2004 on STN
LOCUS (LOC):
GenBank ACC. NO. (GBN):
GenBank VERSION (VER):
                                             BC058617
                                                                   GenBank (R)
                                            BC058617
                                             BC058617.1 GI:35193185
CAS REGISTRY NO. (RN):
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
                                             mRNA; linear
DIVISION CODE (CI):
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DATE (DATE):
                                             30 Jun 2004
DEFINITION (DEF):
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                                                                                                                             mRNA (cDNA
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                                             clone MGC:64741 IMAGE:5695418), complete cds.
KEYWORDS (ST):
                                            MGC
SOURCE:
                                            Mus musculus (house mouse)
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                                            Mus musculus
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                                             Euteleostomi; Mammalia; Eutheria; Rodentia;
                                             Sciurognathi; Muridae; Murinae; Mus
COMMENT:
         Contact: MGC help desk
         Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Dr. Jim Lin, University of Iowa
         cDNA Library Preparation: M. Bento Soares, University of Iowa cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: University of Iowa, Dr. M. Bento Soares and Dr.
         Thomas L. Casavant.
         Web site: http://genome.uiowa.edu
        Contact: bento-soares@uiowa.edu; tom-casavant@uiowa.edu
Bonaldo,M.F., Akabogu,I., Bair,T., Bair,J., Crouch,K., Davis,A.,
Fishler,K., Keppel,C., Kucaba,T., Lebeck,M., Melo,A., Schaefer,K.,
Scheetz,T., Smith,C., Snir,E., Tack,D., Trout,K., Walters,J.,
Casavant,T., Soares,M.B.
         Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Series: Plate: Row: Column: 0
         This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 24475899.
REFERÊNCE:
                                                  (bases 1 to 3135)
                                            Strausberg, R.L.; Feingold, E.A.; Grouse, L.H.;
Derge, J.G.; Klausner, R.D.; Collins, F.S.; Wagner, L.;
Shenmen, C.M.; Schuler, G.D.; Altschul, S.F.; Zeeberg, B.;
Buetow, K.H.; Schaefer, C.F.; Bhat, N.K.; Hopkins, R.F.;
     AUTHOR (AU):
                                             Jordan,H.; Moore,T.; Max,S.I.; Wang,J.; Hsieh,F.;
                                             Diatchenko, L.; Marusina, K.; Farmer, A.A.; Rubin, G.M.;
                                            Hong, L.; Stapleton, M.; Soares, M.B.; Bonaldo, M.F.; Casavant, T.L.; Scheetz, T.E.; Brownstein, M.J.; Usdin, T.B.; Toshiyuki, S.; Carninci, P.; Prange, C.; Raha, S.S.; Loquellano, N.A.; Peters, G.J.; Abramson, R.D.; Mullahy, S.J.; Bosak, S.A.; McEwan, P.J.; McKernan, K.J.; Malek, J.A.; Gunaratne, P.H.; Richards, S.; Worley, K.C.; Hale S.; Garcia A.M.; Cay I. J.; Hulyk S.W.;
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Hale,S.; Garcia,A.M.; Gay,L.J.; Hulyk,S.W.;

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Gibbs,R.A.; Fahey,J.; Helton,E.; Ketteman,M.; Madan,A.;
                                           Rodrigues, S.; Sanchez, A.; Whiting, M.; Madan, A.; Young, A.C.; Shevchenko, Y.; Bouffard, G.G.; Blakesley, R.W.; Touchman, J.W.; Green, E.D.; Dickson, M.C.; Rodriguez, A.C.; Grimwood, J.; Schmutz, J.; Myers, R.M.; Butterfield, Y.S.; Krzywinski, M.I.; Skalska, U.; Smailus, D.E.; Schnerch, A.; Schein, J.E.; Jones, S.J.; Marra, M.A.
     TITLE (TI):
                                           Generation and initial analysis of more than 15,000
                                           full-length human and mouse cDNA sequences
     JOURNAL (SO):
                                           Proc. Natl. Acad. Sci. U.S.A., 99 (26), 16899-16903
                                            (2002)
     OTHER SOURCE (OS):
                                           CA 138:67676
REFERENCE:
                                                 (bases 1 to 3135)
     AUTHOR (AU): TITLE (TI):
                                           Strausberg, R.
                                           Direct Submission
                                           Submitted (22-SEP-2003) National Institutes of Health,
     JOURNAL (SO):
                                           Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA
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        ANSWER 144 OF 159
L7
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COMMENT:
        Contact: MGC help desk
        Email: cgapbs-r@mail.nih.gov
        Tissue Procurement: Dr. Jim Lin, University of Iowa cDNA Library Preparation: M. Bento Soares, University of Iowa cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
        DNA Sequencing by: Sequencing Group at the Stanford Human Genome
        Center, Stanford University School of Medicine, Stanford, CA 94305
        Web site:
                                    http://www-shgc.stanford.edu
                          (Dickson, Mark) mcd@paxil.stanford.edu
        Contact:
        Dickson, M., Schmutz, J., Grimwood, J., Rodriquez, A., and Myers,
        Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
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This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 24475899.
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(bases 1 to 3039)

Strausberg, R.L.; Feingold, E.A.; Grouse, L.H.;

REFERENCE:

AUTHOR (AU):

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Shenmen, C.M.; Schuler, G.D.; Altschul, S.F.; Žeeberg, B.; Buetow, K.H.; Schaefer, C.F.; Bhat, N.K.; Hopkins, R.F.;
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                                   Gibbs, R.A.; Fahey, J.; Helton, E.; Ketteman, M.; Madan, A.; Rodrigues, S.; Sanchez, A.; Whiting, M.; Madan, A.;
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                                    (2002)
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                                   CA 138:67676
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                                   Strausberg, R. Direct Submission
    AUTHOR (AU):
    TITLE (TI):
    JOURNAL (SO):
                                   Submitted (01-MAY-2003) National Institutes of Health,
                                   Mammalian Gene Collection (MGC), Cancer Genomics
                                   Office, National Cancer Institute, 31 Center Drive,
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L7
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   AUTHOR (AU):
TITLE (TI):
JOURNAL (SO):
                        Alitalo, K.; Kubo, H.
                       Ephrin-tie receptor materials and methods
Patent: WO 03004529-A 19 16-JAN-2003; Licentia Ltd.
                        (FI)
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L7 ANSWER 146 OF 159 GENBANK.RTM. LOCUS (LOC): AX671062 GenBank (R) GenBank ACC. NO. (GBN): AX671062 GenBank VERSION (VER): AX671062 CAS REGISTRY NO. (RN): 504048-0 SEQUENCE LENGTH (SQL): 2987 AX671062.1 GI:29329527 504048-06-0 MOLECULE TYPE (CI): DNA; linear DIVISION CODE (CI): Patent DATE (DATE): 27 Mar 2003 DEFINITION (DEF): Sequence 17 from Patent W003004529. SOURCE: Homo sapiens (human) ORGANISM (ORGN): Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo NUCLEIC ACID COUNT (NA): 529 a 886 c 864 q 708 t REFERENCE: AUTHOR (AU): Alitalo, K.; Kubo, H. Ephrin-tie receptor materials and methods TITLE (TI):

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JOURNAL (SO):

Location Feature Key Qualifier

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L7 ANSWER 147 OF 159 GENBANK.RTM. COPYRIGHT 2004 on STN

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GenBank VERSION (VER): BC042944.2 GI:34193522

CAS REGISTRY NO. (RN): 495664-88-5

SEQUENCE LENGTH (SQL): 2338

MOLECULE TYPE (CI): mRNA; linear

DIVISION CODE (CI): Primates
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DATE (DATE): 30 Jun 2004
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SOURCE: Homo sapiens (human)
ORGANISM (ORGN): Homo sapiens

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Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
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  COMMENT:
               On Aug 25, 2003 this sequence version replaced gi:27696534. Contact: MGC help desk
              Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI) & Shiraki
               Toshiyuki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
              DNA Sequencing by: Genome Sequence Centre, BC Cancer Agency, Vancouver, BC, Canada
             info@bcgsc.bc.ca
Steve Jones, Sarah Barber, Mabel Brown-John, Yaron Butterfield,
Andy Chan, Steve S. Chand, William Chow, Alison Cloutier, Ruth
Featherstone, Malachi Griffith, Obi Griffith, Ran Guin, Nancy Liao,
Kim MacDonald, Amara Masson, Mike R. Mayo, Josh Moran, Ryan Morin,
Teika Olson, Diana Palmquist, Anca Petrescu, Anna Liisa Prahbu,
Parvaneh Saeedi, JR Santos, Angelique Schnerch, Ursula Skalska,
Duane Smailus, Jeff Stott, Miranda Tsai, George Yang, Jacquie
Schein, Asim Siddiqui, Rob Holt, Marco Marra.
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Series: IRAK Plate: 76 Row: a Column: 9
This clone was selected for full length sequencing because it
passed the following selection criteria: matched mRNA qi: 27894382.
               info@bcqsc.bc.ca
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Casavant,T.L.; Scheetz,T.E.; Brownstein,M.J.;
Usdin.T.B.: Toshivuki,S.; Carninci,P.; Prange,C.;
         AUTHOR (AU):
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                                                                Skalska, U.; Smailus, D.E.; Schnerch, A.; Schein, J.E.; Jones, S.J.; Marra, M.A. Generation and initial analysis of more than 15,000
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Proc. Natl. Acad. Sci. U.S.A., 99 (26), 16899-16903
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                                                                Mammalian Gene Collection (MGC), Cancer Genomics
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L7 ANSWER 148 OF 159 GENBANK.RTM. COPYRIGHT 2004 on STN

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CAS REGISTRY NO. (RN): 495565-98-5
SEQUENCE LENGTH (SQL): 3394
MOLECULE TYPE (CI): DNA; linear
DIVISION CODE (CI): Patent
DATE (DATE): 8 Jan 2003
DEFINITION (DEF): Sequence 227 from Patent

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Homo sapiens (human)

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    AUTHOR (AU):
                                 Garcia,T.; roman Roman,S.; Baron,R.; Call,K.;
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    TITLE (TI):
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        Division of Experimental Animal Research in Riken contributed to
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Please visit our web site for further details.
        URL:http://genome.gsc.riken.jp/
URL:http://fantom.gsc.riken.jp/.
REFERENCE:
     AUTHOR (AU):
                                     Carninci, P.; Hayashizaki, Y.
     TITLE (TI):
                                     High-efficiency full-length cDNA cloning
     JOURNAL (SO):
                                    Meth. Enzymol., 303, 19-44 (1999)
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REFERENCE:
                                    Carninci, P.; Shibata, Y.; Hayatsu, N.; Sugahara, Y.; Shibata, K.; Itoh, M.; Konno, H.; Okazaki, Y.; Muramatsu, M.; Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid
     AUTHOR (AU):
     TITLE (TI):
                                    discovery of new genes
Genome Res., 10 (10), 1617-1630 (2000)
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    AUTHOR (AU):
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    TITLE (TI):
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    AUTHOR (AU):
                                    The RIKEN Genome Exploration Research Group Phase II
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    TITLE (TI):
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CA 134:203311
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    AUTHOR (AU):
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    TITLE (TI):
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Adachi, J.; Aizawa, K.; Akimura, T.; Arakawa, T.; Bono, H.; Carninci, P.; Fukuda, S.; Furuno, M.; Hanagaki, T.; Hara, A.; Hashizume, W.; Hayashida, K.; Hayatsu, N.; Hiramoto, K.; Hiraoka, T.; Hirozane, T.; Hori, F.; Imotani, K.; Ishii, Y.; Itoh, M.; Kagawa, I.; Kasukawa, T.; Katoh, H.; Kawai, J.; Kojima, Y.; Kondo, S.; Konno, H.; Kouda, M.; Koya, S.; Kurihara, C.; Matsuyama, T.; Miyazaki, A.; Murata, M.; Nakamura, M.; Nishi, K.;
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                                                     Institute of Physical and Chemical Research (RIKEN)
                                                    Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.jp, URL:http://genome.gsc.riken.jp/, Tel:81-45-503-9222, Fax:81-45-503-9216)
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          ANSWER 150 OF 159
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LOCUS (LOC):
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GenBank VERSION (VER):
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MOLECULE TYPE (CI):
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DIVISION CODE (CI):
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DATE (DATE):
                                            4 Feb 2002
DEFINITION (DEF):
                                            Homo sapiens, Similar to
                                                                                             ***ephrin***
                                                                                                                              ***B3***
                                            clone IMAGE: 4814898, mRNA, partial cds.
SOURCE:
                                            human.
  ORGANISM (ORGN):
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                                            Hominidae; Homo
NUCLEIC ACID COUNT (NA): 532 a
                                                             788 c
                                                                           703 q
                                                                                          650 t
COMMENT:
         Contact: MGC help desk
         Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI) &
         Toshiyuki and Piero Carninci (RIKEN)

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Sequencing Group at the Stanford Human Genome

Center, Stanford University School of Medicine, Stanford, CA 94305
                                                                                                                    Shiraki
         Web site:
                                      http://www-shgc.stanford.edu
                            (Dickson, Mark) mcd@paxil.stanford.edu
         Contact:
         Dickson, M., Schmutz, J., Grimwood, J., Rodriquez, A., and Myers,
         Clone distribution: MGC clone distribution information can be found
         through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Series: IRAK Plate: 32 Row: 1 Column: 19.
REFERENCE:
                                                  (bases 1_to 2673)
     AUTHOR (AU):
                                            Strausberg, R.
     TITLE (TI):
                                           Direct Submission
     JOURNAL (SO):
                                           Submitted (01-FEB-2002) National Institutes of Health,
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Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA

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ANSWER 151 OF 159
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 LOCUS (LOC): BI313373
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GenBank VERSION (VER): BI313373
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  DATE (DATE):
                                       20 Jul 2001
                                      dah92f01.y1 NICHD XGC Emb4 Xenopus laevis cDNA clone IMAGE:4957560 5' similar to TR:Q9PT69 Q9PT69 ***EPHRIN*** - ***B3*** PRECURSOR.;, mRNA
  DEFINITION (DEF):
                                       sequence.
  SOURCE:
                                      African clawed frog.
   ORGANISM (ORGN):
                                      Xenopus laevis
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                                      Euteleostomi; Amphibia; Batrachia; Anura;
                                      Mesobatrachia; Pipoidea; Pipidae; Xenopodinae; Xenopus
 NUCLEIC ACID COUNT (NA): 118 a
                                                               92 g
                                                   152 c
                                                                            94 t
 COMMENT:
         Other ESTs: dah92f01.x1
         Contact: Robert Strausberg, Ph.D.
         Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Martha Rebbert, Steven L. Klein, Ph.D.
          cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: Xenopus clones from this library are available
         through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov
Seq primer: -40RP from Gibco
         High quality sequence stop: 434.
 REFERENCE:
                                           (bases 1 to 456)
                                     NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
     AUTHOR (AU): TITLE (TI):
                                     National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
     JOURNAL (SO):
                                     Unpublished (1997)
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                                                              unidirectionally. Primer: Oligo
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Constructed by Life Technologies.
Note: This is a Xenopus Gene
Collection (XGC) library."
SEQUENCE (SEQ):
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LOCUS (LOC):
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GenBank ACC. NO. (GBN): AF375227
GenBank VERSION (VER):
                                    AF375227.1 GI:14495333
CAS REGISTRY NO. (RN):
                                     342875-57-4
SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
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DATE (DATE):
                                         20 Jun 2001
  DEFINITION (DEF):
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                                                               ***ephrin***
                                                                                         ***B3***
                                                                                                          mRNA,
                                         complete cds.
  SOURCE:
                                         zebrafish.
    ORGANISM (ORGN):
                                         Danio rerio
                                        Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio 231 a 293 c 256 g 216 t
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                                              (bases 1 to 996)
       AUTHOR (AU):
                                        Chan, J.; Mably, J.D.; Serluca, F.C.; Chen, J.N.; Goldstein, N.B.; Thomas, M.C.; Cleary, J.A.; Brennan, C.; Fishman, M.C.; Roberts, T.M.
                                        Morphogenesis of prechordal plate and notochord requires intact Eph/ephrin B signaling Dev. Biol., 234 (2), 470-482 (2001)
       TITLE (TI):
       JOURNAL (SO):
       OTHER SOURCE (OS):
                                        CA 135:164797
  REFERENCE:
                                             (bases 1 to 996)
      AUTHOR (AU):
                                        Chan, J.; Mably, J.D.; Serluca, F.C.; Chen, J.-N.;
                                        Goldstein, N.B.; Thomas, M.C.; Cleary, J.A.; Brennan, C.; Fishman, M.C.; Roberts, T.M.
      TITLE (TI):
                                        Direct Submission
      JOURNAL (SO):
                                        Submitted (01-MAY-2001) Department of Cancer Biology, Dana-Farber Cancer Institute, 1 Jimmy Fund Way, Boston,
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        ANSWER 153 OF 159
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LOCUS (LOC):
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GenBank VERSION (VER):
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MOLECULE TYPE (CI):
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                                                                                     30 Jan 2004
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                                                                                    moderately similar to ***EPHRIN***
                                                                                    PRECURSOR.
   KEYWORDS (ST):
                                                                                    oligo capping; fis (full insert sequence)
   SOURCE:
                                                                                    Homo sapiens (human)
      ORGANISM (ORGN):
                                                                                    Homo sapiens
                                                                                    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
                                                                                    Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini;
                                                                                    Hominidae; Homo
   COMMENT:
                  NEDO human cDNA sequencing project supported by Ministry of Economy, Trade and Industry of Japan; cDNA full insert sequencing: Research Association for Biotechnology; cDNA library construction, 5'- & 3'-end one pass sequencing and clone selection: Helix Research Institute (supported by Japan Key Technology Center etc.) and Department of Virology, Institute of Medical Science, University of Tokyo.
  REFERENCE:
            AUTHOR (AU):
                                                                                   Ota, T.; Suzuki, Y.; Nishikawa, T.; Otsuki, T.;
                                                                                 Ota, T.; Suzuki, Y.; Nishikawa, T.; Otsuki, T.; Sugiyama, T.; Irie, R.; Wakamatsu, A.; Hayashi, K.; Sato, H.; Nagai, K.; Kimura, K.; Makita, H.; Sekine, M.; Obayashi, M.; Nishi, T.; Shibahara, T.; Tanaka, T.; Ishii, S.; Yamamoto, J.; Saito, K.; Kawai, Y.; Isono, Y.; Nakamura, Y.; Nagahari, K.; Murakami, K.; Yasuda, T.; Iwayanagi, T.; Wagatsuma, M.; Shiratori, A.; Sudo, H.; Hosoiri, T.; Kaku, Y.; Kodaira, H.; Kondo, H.; Sugawara, M.; Takahashi, M.; Kanda, K.; Yokoi, T.; Furuya, T.; Kikkawa, E.; Omura, Y.; Abe, K.; Kamihara, K.; Katsuta, N.;
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Kikkawa, E.; Omura, Y.; Abe, K.; Kamihara, K.; Katsuta, N.;
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Yosida, M.; Hotuta, T.; Kusano, J.; Kanehori, K.;
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Togiya, S.; Komai, F.; Hara, R.; Takeuchi, K.; Arita, M.;
Imose, N.; Musashino, K.; Yuuki, H.; Oshima, A.; Sasaki, N.;
Aotsuka, S.; Yoshikawa, Y.; Matsunawa, H.; Ichihara, T.;
Shiohata, N.; Sano, S.; Moriya, S.; Momiyama, H.; Satoh, N.;
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                                                                                 Sugano, S.
           TITLE (TI):
                                                                                 Complete sequencing and characterization of 21,243
                                                                                 full-length human cDNAs
           JOURNAL (SO):
                                                                                 Nat. Genet., 36 (1), 40-45 (2004)
          OTHER SOURCE (OS):
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                                                                                Isogai, T.; Ota, T.; Hayashi, K.; Sugiyama, T.; Otsuki, T.; Suzuki, Y.; Nishikawa, T.; Nagai, K.; Sugano, S.; Shiratori, A.; Sudo, H.; Wagatsuma, M.; Hosoiri, T.; Kaku, Y.; Kodaira, H.; Kondo, H.; Sugawara, M.; Takahashi, M.; Chiba, Y.; Ishida, S.; Murakawa, K.; Ono, Y.; Takiguchi, S.; Watanabe, S.; Kimura, K.; Murakami, K.; Ishii S.; Kawai V.; Saito K.; Vamamoto J.;
          AUTHOR (AU):
                                                                                 Ishii,S.; Kawai,Y.; Saito,K.; Yamamoto,J.;
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          TITLE (TI):
                                                                                NEDO human cDNA sequencing project
          JOURNAL (SO):
                                                                                Unpublished
REFERENCE:
                                                                                            (bases 1 to 2153)
          AUTHOR (AU):
                                                                                Isogai,T.; Otsuki,T.
Direct Submission
          TITLE (TI):
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Institute, Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan (E-mail:genomics@hri.co.jp, Tel:81-438-52-3975, Fax:81-438-52-3986)
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L7
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                                                      COPYRIGHT 2004 on STN
LOCUS (LOC):
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GenBank VERSION (VER):
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SEQUENCE LENGTH (SQL):
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DATE (DATE):
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DEFINITION (DEF):
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                                to TR:Q9PT69 Q9PT69
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                                PRECURSOR. ;, mRNA sequence.
SOURCE:
                                western clawed frog.
 ORGANISM (ORGN):
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                                Euteleostomi; Amphibia; Batrachia; Anura;
Mesobatrachia; Pipoidea; Pipidae; Xenopodinae; Silurana
137 a 129 c 124 g 166 t
NUCLEIC ACID COUNT (NA): 137 a 129 c
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Other ESTs: dad19b07.y1
              Contact: Sandy Clifton, Ph.D. WashU Xenopus EST project, 1999
              Washington University School of Medicine
              4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800
Fax: 314 286 1810
              Email: est@watson.wustl.edu
              Library constructed by A. Zorn and J. Mason (Wellcome/CRC Institute). DNA Sequencing by: Washington University Genome Sequencing
              Center
                Clone distribution: Xenopus clones from this library are available
              through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov
              Seq primer: -40UP from Gibco
              High quality sequence stop: 471.
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                                                         Clifton, S.; Johnson, S.L.; Blumberg, B.; Song, J.; Hillier, L.; Pape, D., Martin, J.; Wylie, T.; Underwood, K.; Theising, B.; Bowers, Y.; Person, B.M; Gibbons, M.; Harvey, N.; Ritter, E.; Jackson, Y.; McCann, R.; Waterston, R.; Wilson, R.
         AUTHOR (AU):
                                                         WashU Xenopus EST project, 1999
Unpublished (1999)
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         JOURNAL (SO):
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COMMENT:
            Contact: Yoshihide Hayashizaki
           Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan Tel: 81-45-503-9222
           Fax: 81-45-503-9216
           Email: genome-res@gsc.riken.go.jp,
           URL: http://genome.gsc.riken.go.jp/
Carninci,P., Nishiyama,Y., Westover,A., Itoh,M., Nagaoka,S., Sasaki,N., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
Thermostabilization and thermoactivation of thermolabile enzymes by
           trehalose and its application for the synthesis of full length cDNA. Proc. Natl. Acad. Sci. U.S.A. 95 (2), 520-524 (1998) Itoh, M., Kitsunai, T., Akiyama, J., Shibata, K., Izawa, M., Kawai, J., Tomaru, Y., Carninci, P., Shibata, Y., Ozawa, Y., Muramatsu, M., Okazaki V. and Havashizaki V.
           ,Y. and Hayashizaki,Y.
           Automated filtration-based high-throughput plasmid preparation system. Genome Res. 9 (5), 463-470 (1999)
            Carninci, P. and Hayashizaki, Y. High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
           19 - 44 (1999)
            Please visit our web site (http://genome.rtc.riken.go.jp) for
           further details.
 REFERENCE:
                                               1 (bases 1 to 301)
Konno,H.; Aizawa,K.; Akahira,S.; Akiyama,J.;
Arakawa,T.; Carninci,P.; Endo,T.; Fukuda,S.;
Fukunishi,Y.; Hara,A.; Hayatsu,N.; Hirozane,T.;
Hori,F.; Ishii,Y.; Ishikawa,J.; Ishikawa,T.; Itoh,M.;
Izawa,M.; Kadota,K.; Kagawa,I.; Kai,C.; Kawai,J.;
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Mizuno,Y.; Nakamura,M.; Oda,H.; Okazaki,Y., Ono,T.y;
Owa,C.; Saito,H.; Sakai,C.; Sato,K.; Shibata,K.;
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                                                                               Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken
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thermo-activated reverse

transcriptase and subsequently enriched for full-length by

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17 Apr 2000
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DATE (DATE):
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                                  Comparative analysis of embryonic gene expression defines potential interaction sites for Xenopus EphB4
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                                  Brandli, A.W.
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        TITLE (TI):
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***Ephrin*** - ***B3***, a ligand for the receptor EphB3, expressed at the midline of the developing neural tube
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        Effects of pre- and postnatal methylmercury exposure on expression of EPHS
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AU
        Wilson D T; Reuhl K R; Zhou R
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       Ricard, J.; Salinas, J.A.; Liebl, D.J.
University of Miami, School of Medicine; URL: www.med.miami.edu.
Meeting Info.: 000 7201: 2004 Miami Nature Biotechnology Winter Symposium (0007201). Miami Beach, FL (USA). 31 Jan-4 Feb 2004. University of Miami.
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